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FORM PTO-1390 (Modified)
(REV 10-95)

U.S. DEPARTMENT OF COMMERCE PATENT AND TRADEMARK OFFICE

ATTORNEY'S DOCKET NUMBER

TRANSMITTAL LETTER TO THE UNITED STATES

DESIGNATED/ELECTED OFFICE (DO/EO/US)

CONCERNING A FILING UNDER 35 U.S.C. 371

18-971-0 PCT

U.S. APPLICATION NO. (IF KNOWN, SEE 37 CFR 1.5)

08/809723

INTERNATIONAL APPLICATION NO.

PCT/JP95/01983

INTERNATIONAL FILING DATE

29 September 1995

PRIORITY DATE CLAIMED

07 October 1994, earliest

TITLE OF INVENTION

NEW COMPOUND

APPLICANT(S) FOR DO/EO/US

Hidenori OHKI, et al

Applicant herewith submits to the United States Designated/Elected Office (DO/EO/US) the following items and other information:

1. ☒ This is a **FIRST** submission of items concerning a filing under 35 U.S.C. 371.
2. ☐ This is a **SECOND** or **SUBSEQUENT** submission of items concerning a filing under 35 U.S.C. 371.
3. ☒ This express request to begin national examination procedures (35 U.S.C. 371(f)) at any time rather than delay examination until the expiration of the applicable time limit set in 35 U.S.C. 371(b) and PCT Articles 22 and 39(1).
4. ☒ A proper Demand for International Preliminary Examination was made by the 19th month from the earliest claimed priority date.
5. ☒ A copy of the International Application as filed (35 U.S.C. 371 (c) (2))
 - a. ☐ is transmitted herewith (required only if not transmitted by the International Bureau).
 - b. ☒ has been transmitted by the International Bureau.
 - c. ☐ is not required, as the application was filed in the United States Receiving Office (RO/US).
6. ☒ A translation of the International Application into English (35 U.S.C. 371(c)(2)).
7. ☒ A copy of the International Search Report (PCT/ISA/210).
8. ☒ Amendments to the claims of the International Application under PCT Article 19 (35 U.S.C. 371 (c)(3))
 - a. ☐ are transmitted herewith (required only if not transmitted by the International Bureau).
 - b. ☐ have been transmitted by the International Bureau.
 - c. ☐ have not been made; however, the time limit for making such amendments has NOT expired.
 - d. ☒ have not been made and will not be made.
9. ☐ A translation of the amendments to the claims under PCT Article 19 (35 U.S.C. 371(c)(3)).
10. ☐ An oath or declaration of the inventor(s) (35 U.S.C. 371 (c)(4)).
11. ☒ A copy of the International Preliminary Examination Report (PCT/IPEA/409).
12. ☐ A translation of the annexes to the International Preliminary Examination Report under PCT Article 36 (35 U.S.C. 371 (c)(5)).

Items 13 to 18 below concern document(s) or information included:

13. ☐ An Information Disclosure Statement under 37 CFR 1.97 and 1.98.
14. ☐ An assignment document for recording. A separate cover sheet in compliance with 37 CFR 3.28 and 3.31 is included.
15. ☒ A **FIRST** preliminary amendment.
A **SECOND** or **SUBSEQUENT** preliminary amendment.
16. ☐ A substitute specification.
17. ☐ A change of power of attorney and/or address letter.
18. ☒ Other items or information:

Notice of Priority

PCT/IB/304

PCT/IB/308

Request for Consideration of Documents Cited in International Search Report

U.S. APPLICATION NO. (IF KNOWN, SEE 37 CFR 1.5)

INTERNATIONAL APPLICATION NO.

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19. The following fees are submitted:

BASIC NATIONAL FEE (37 CFR 1.492 (a) (1) - (5)) :

Search Report has been prepared by the EPO or JPO	\$910.00
International preliminary examination fee paid to USPTO (37 CFR 1.482)	\$700.00
No international preliminary examination fee paid to USPTO (37 CFR 1.482) but international search fee paid to USPTO (37 CFR 1.445(a)(2))	\$770.00
Neither international preliminary examination fee (37 CFR 1.482) nor international search fee (37 CFR 1.445(a)(2) paid to USPTO	\$1,040.00
International preliminary examination fee paid to USPTO (37 CFR 1.482) and all claims satisfied provisions of PCT Article 33(2)-(4)	\$96.00

ENTER APPROPRIATE BASIC FEE AMOUNT =

\$910.00

Surcharge of \$130.00 for furnishing the oath or declaration later than ☐ 20 ☒ 30 months from the earliest claimed priority date (37 CFR 1.492 (e)).

\$130.00

CLAIMS	NUMBER FILED	NUMBER EXTRA	RATE
Total claims	19 - 20 =	0	x \$22.00
Independent claims	2 - 3 =	0	x \$80.00

\$0.00

\$0.00

Multiple Dependent Claims (check if applicable). ☐

\$0.00

TOTAL OF ABOVE CALCULATIONS =

\$1,040.00

Reduction of 1/2 for filing by small entity, if applicable. Verified Small Entity Statement must also be filed (Note 37 CFR 1.9, 1.27, 1.28) (check if applicable). ☐

\$0.00

SUBTOTAL =

\$1,040.00

Processing fee of \$130.00 for furnishing the English translation later than ☐ 20 ☐ 30 months from the earliest claimed priority date (37 CFR 1.492 (f)). +

\$0.00

TOTAL NATIONAL FEE =

\$1,040.00

Fee for recording the enclosed assignment (37 CFR 1.21(h)). The assignment must be accompanied by an appropriate cover sheet (37 CFR 3.28, 3.31) (check if applicable). ☐

\$0.00

TOTAL FEES ENCLOSED =

\$1,040.00

Amount to be:
refunded

\$

charged

\$

☒ A check in the amount of \$1,040.00 to cover the above fees is enclosed.☐ Please charge my Deposit Account No. _____ in the amount of _____ to cover the above fees.
A duplicate copy of this sheet is enclosed.☒ The Commissioner is hereby authorized to charge any fees which may be required, or credit any overpayment to Deposit Account No. 15-0030 A duplicate copy of this sheet is enclosed.**NOTE: Where an appropriate time limit under 37 CFR 1.494 or 1.495 has not been met, a petition to revive (37 CFR 1.137(a) or (b)) must be filed and granted to restore the application to pending status.**

SEND ALL CORRESPONDENCE TO:

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NAME

24,618

REGISTRATION NUMBER

DATE

April 7, 1997

DESCRIPTION

NEW COMPOUND

5 TECHNICAL FIELD

The present invention relates to new polypeptide compound and a pharmaceutically acceptable salt thereof which are useful as a medicament.

10 BACKGROUND ART

In U.S. Pat. No. 5,376,634, there are disclosed the polypeptide compound and a pharmaceutically acceptable salt thereof, which have antimicrobial activities (especially antifungal activity).

15

DISCLOSURE OF INVENTION

The present invention relates to new polypeptide compound and a pharmaceutically acceptable salt thereof.

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More particularly, it relates to new polypeptide compound and a pharmaceutically acceptable salt thereof, which have antimicrobial activities [especially, antifungal activities, in which the fungi may include Aspergillus, Cryptococcus, Candida, Mucor, Actinomyces, Histoplasma, Dermatophyte, Malassezia, Fusarium and the like.], inhibitory activity on β -1,3-glucan synthase, and further which are expected to be useful for the prophylactic and/or therapeutic treatment of Pneumocystis carinii infection (e.g. Pneumocystis carinii pneumonia) in a human being or an animal, to a process for preparation thereof, to a pharmaceutical composition comprising the same, and to a method for the prophylactic and/or therapeutic treatment of infectious diseases including Pneumocystis carinii infection (e.g. Pneumocystis carinii pneumonia) in a human being or an animal.

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lower alkanoyl substituted with
unsaturated condensed heterocyclic group
containing 1 to 3 sulfur atom(s) which may
have one or more suitable substituent(s);
lower alkanoyl substituted with

aryl substituted with aryl having lower alkyl;

aryl substituted with aryl having higher alkyl;

5 aryloxy(lower)alkanoyl which may have one or more suitable substituent(s);

ar(lower)alkoxy(lower)alkanoyl which may have one or more suitable substituent(s);

10 arylamino(lower)alkanoyl which may have one or more suitable substituent(s);

lower alkanoyl substituted with pyrazolyl which has lower alkyl and aryl having higher alkoxy;

15 lower alkoxy(higher)alkanoyl, in which higher alkanoyl may have one or more suitable substituent(s);

20 aryl substituted with aryl having heterocyclicoxy, in which heterocyclicoxy may have one or more suitable substituent(s);

aryl substituted with cyclo(lower)alkyl having lower alkyl;

indolylcarbonyl having higher alkyl;

naphthoyl having lower alkyl;

25 naphthoyl having higher alkyl;

naphthoyl having lower alkoxy(higher)alkoxy;

aryl substituted with aryl having lower alkoxy(lower)alkoxy(higher)alkoxy;

30 aryl substituted with aryl having lower alkoxy(lower)alkoxy;

aryl substituted with aryl which has aryl having lower alkoxy;

35 aryl substituted with aryl which has aryl having lower alkoxy(lower)alkoxy;

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aroyl substituted with aryl having
heterocyclicoxy(higher)alkoxy;

aroyl substituted with aryl having
aryloxy(lower)alkoxy;

5 aroyl substituted with aryl having
heterocycliccarbonyl(higher)alkoxy;

lower alkanoyl substituted with oxazolyl
which has aryl having higher alkoxy;

10 lower alkanoyl substituted with furyl
which has aryl substituted with aryl having
lower alkoxy;

lower alkanoyl substituted with triazolyl
which has oxo and aryl having higher alkyl;

15 higher alkanoyl having hydroxy;

higher alkanoyl having ar(lower)alkyl and
hydroxy;

3-methyl-tridecenoyl; or

20 (C₂-C₆)alkanoyl substituted with aryl
having higher alkoxy, in which (C₂-
C₆)alkanoyl may have amino or protected
amino.

25 The new polypeptide compound [I] and a
pharmaceutically acceptable salt thereof can be prepared
by the process as illustrated in the following reaction
scheme or can be prepared by elimination reaction of amino
protective group in R¹.

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wherein R¹ is as defined above.

Suitable pharmaceutically acceptable salts of the object polypeptide compound [I] are conventional non-toxic salts and may include a salt with a base or an acid addition salt such as a salt with an inorganic base, for example, an alkali metal salt (e.g., sodium salt, potassium salt, etc.), an alkaline earth metal salt (e.g., calcium salt, magnesium salt, etc.), an ammonium salt; a salt with an organic base, for example, an organic amine salt (e.g., triethylamine salt, pyridine salt, picoline salt, ethanolamine salt, triethanolamine salt, dicyclohexylamine salt, N,N'-dibenzylethylenediamine salt, etc.); an inorganic acid addition salt (e.g., hydrochloride, hydrobromide, sulfate, phosphate, etc.); an organic carboxylic sulfonic acid addition salt (e.g., formate, acetate, trifluoroacetate, maleate, tartrate, fumarate, methanesulfonate, benzenesulfonate, toluenesulfonate, etc.); a salt with a basic or acidic amino acid (e.g., arginine, aspartic acid, glutamic acid, etc.).

In the above and subsequent descriptions of the present specification, suitable examples and illustration of the various definitions which the present invention intends to include within the scope thereof are explained in detail as follows.

The term "lower" is used to intend a group having 1 to 6 carbon atom(s), unless otherwise provided.

The term "higher" is used to intend a group having 7 to 20 carbon atoms, unless otherwise provided.

Suitable example of "one or more" may be the number of 1 to 6, in which the preferred one may be the number of 1 to 3.

Suitable example of "lower alkanoyl" may include

straight or branched one such as formyl, acetyl, 2-methylacetyl, 2,2-dimethylacetyl, propionyl, butyryl, isobutyryl, pentanoyl, 2,2-dimethylpropionyl, hexanoyl, and the like.

- 5 Suitable example of "suitable substituent(s)" in the groups such as "lower alkanoyl substituted with unsaturated 6-membered heteromonocyclic group containing at least one nitrogen atom which may have one or more suitable substituent(s)", "lower alkanoyl substituted with 10 1,2,3,4-tetrahydroisoquinoline which may have one or more suitable substituent(s)", etc. may include lower alkoxy as mentioned below, higher alkoxy as mentioned below, lower alkyl as mentioned below, higher alkyl as mentioned below, higher alkoxy(lower)alkyl, lower alkoxycarbonyl, oxo, aryl 15 which may have one or more lower alkoxy, aryl which may have one or more higher alkoxy, aryl which may have one or more lower alkyl, aryl which may have one or more higher alkyl, aryl substituted with aryl which may have one or more lower alkoxy, aryl substituted with aryl which may have one or more higher alkoxy, aryl substituted with aryl which may 20 have one or more lower alkyl, aryl substituted with aryl which may have one or more higher alkyl, aroyl which may have one or more lower alkoxy, aroyl which may have one or more higher alkoxy, aroyl which may have one or more lower alkyl, aroyl which may have one or more higher alkyl, heterocyclic group which may have one or more lower alkoxy, heterocyclic group which may have one or more higher alkoxy, aryl having 25 heterocyclic(higher)alkoxy, heterocyclic group which may have aryl having higher alkoxy, heterocyclic group which may have aryl having lower alkoxy(higher)alkoxy, heterocyclic group which may have aryl having lower alkoxy, lower alkoxy(lower)alkyl, halo(lower)alkoxy, lower alkenyloxy, halo(higher)alkoxy, lower 30 alkoxy(higher)alkoxy, aryl which may have one or more

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lower alkoxy(lower)alkoxy, heterocyclic group, aryl which may have one or more lower alkoxy(higher)alkoxy, aryl which may have one or more higher alkenyloxy, cyclo(lower)alkyl which may have aryl, aryl substituted with heterocyclic group which may have lower alkyl and oxo, cyclo(lower)alkyl which may have one or more lower alkyl, aryl which may have cyclo(lower)alkyl, aryl which may have heterocyclic group, and the like.

Suitable example of "lower alkoxy" may include straight or branched one such as methoxy, ethoxy, propoxy, isopropoxy, butoxy, isobutoxy, tert-butoxy, pentyloxy, tert-pentyloxy, neo-pentyloxy, hexyloxy, isohexyloxy and the like,

in which the preferred one may be methoxy, ethoxy, propoxy, butoxy, pentyloxy, hexyloxy and isohexyloxy.

Suitable example of "higher alkoxy" may include straight or branched one such as heptyloxy, octyloxy, 3,5-dimethyloctyloxy, 3,7-dimethyloctyloxy, nonyloxy, decyloxy, undecyloxy, dodecyloxy, tridecyloxy, tetradecyloxy, hexadecyloxy, heptadecyloxy, octadecyloxy, nonadecyloxy, icosyloxy, and the like,

in which the preferred one may be (C₇-C₁₄)alkoxy, and the more preferred one may be heptyloxy and octyloxy.

Suitable example of "lower alkyl" may include straight or branched one having 1 to 6 carbon atom(s), such as methyl, ethyl, propyl, isopropyl, butyl, isobutyl, sec-butyl, tert-butyl, pentyl, tert-pentyl, neo-pentyl, hexyl, isohexyl and the like,

in which the preferred one may be methyl, pentyl, hexyl and isohexyl.

Suitable example of "higher alkyl" may include straight or branched one having 7 to 20 carbon atoms, such as heptyl, octyl, 3,5-dimethyloctyl, 3,7-dimethyloctyl, nonyl, decyl, undecyl, dodecyl, tridecyl, tetradecyl, pentadecyl, hexadecyl, heptadecyl, octadecyl, nonadecyl,

icosyl, and the like,

in which the preferred one may be (C₇-C₁₄)alkyl, and the more preferred one may be heptyl, octyl, nonyl and decyl.

Suitable example of "aryl" and "ar" moiety may
5 include phenyl which may have lower alkyl (e.g., phenyl, mesityl, tolyl, etc.), naphthyl, anthryl, and the like, in which the preferred one may be phenyl and naphthyl.

Suitable example of "aroyl" may include benzoyl, toluoyl, naphthoyl, anthrylcarbonyl, and the like,
10 in which the preferred one may be benzoyl and naphthoyl.

Suitable example of "heterocyclic group" and "heterocyclic" moiety may include

unsaturated 3 to 8-membered (more preferably 5 or 6-
15 membered) heteromonocyclic group containing 1 to 4 nitrogen atom(s), for example, pyrrolyl, pyrrolinyl, imidazolyl, pyrazolyl, pyridyl, dihydropyridyl, pyrimidyl, pyrazinyl, pyridazinyl, triazolyl (e.g., 4H-1,2,4-triazolyl, 1H-1,2,3-triazolyl, 2H-1,2,3-triazolyl, etc.),
20 tetrazolyl (e.g., 1H-tetrazolyl, 2H-tetrazolyl, etc.), etc.;

saturated 3 to 8-membered (more preferably 5 or 6-membered heteromonocyclic group containing 1 to 4 nitrogen atom(s), for example, pyrrolidinyl, imidazolidinyl,
25 piperidyl, piperazinyl, etc.;

unsaturated condensed heterocyclic group containing 1 to 4 nitrogen atom(s), for example, indolyl, isoindolyl, indolinyl, indoliziny, benzimidazolyl, quinolyl, isoquinolyl, indazolyl, benzotriazolyl, etc.;

unsaturated 3 to 8-membered (more preferably 5 or 6-membered) heteromonocyclic group containing 1 to 2 oxygen atom(s) and 1 to 3 nitrogen atom(s), for example, oxazolyl, isoxazolyl, oxadiazolyl (e.g., 1,2,4-oxadiazolyl, 1,3,4-oxadiazolyl, 1,2,5-oxadiazolyl, etc.),
30 etc.;

saturated 3 to 8-membered (more preferably 5 or 6-membered) heteromonocyclic group containing 1 to 2 oxygen atom(s) and 1 to 3 nitrogen atom(s), for example, morpholinyl, sydnonyl, etc.;

5 unsaturated condensed heterocyclic group containing 1 to 2 oxygen atom(s) and 1 to 3 nitrogen atom(s), for example, benzoxazolyl, benzoxadiazolyl, etc.;

10 unsaturated 3 to 8-membered (more preferably 5 or 6-membered) heteromonocyclic group containing 1 to 2 sulfur atom(s) and 1 to 3 nitrogen atom(s), for example, thiazolyl, isothiazolyl, thiadiazolyl (e.g., 1,2,3-thiadiazolyl, 1,2,4-thiadiazolyl, 1,3,4-thiadiazolyl, 1,2,5-thiadiazolyl, etc.), dihydrothiazinyl, etc.;

15 saturated 3 to 8-membered (more preferably 5 or 6-membered) heteromonocyclic group containing 1 to 2 sulfur atom(s) and 1 to 3 nitrogen atom(s), for example, thiazolidinyl, etc.;

20 unsaturated 3 to 8-membered (more preferably 5 or 6-membered) heteromonocyclic group containing 1 to 2 sulfur atom(s), for example, thienyl, dihydrodithiynyl, dihydrodithionyl, etc.;

25 unsaturated condensed heterocyclic group containing 1 to 2 sulfur atom(s) and 1 to 3 nitrogen atom(s), for example, benzothiazolyl, benzothiadiazolyl, etc.;

30 unsaturated 3 to 8-membered (more preferably 5 or 6-membered) heteromonocyclic group containing an oxygen atom, for example, furyl, etc.;

35 saturated 3 to 8-membered (more preferably 5 or 6-membered) heteromonocyclic group containing an oxygen atom, for example, tetrahydrofuran, tetrahydropyran, etc.;

 unsaturated 3 to 8-membered (more preferably 5 or 6-membered) heteromonocyclic group containing an oxygen atom and 1 to 2 sulfur atom(s), for example, dihydrooxathiinyl, etc.;

 unsaturated condensed heterocyclic group containing 1

to 2 sulfur atom(s), for example, benzothienyl, benzodithiynyl, etc.;

unsaturated condensed heterocyclic group containing an oxygen atom and 1 to 2 sulfur atom(s), for example,
5 benzoxathiynyl, etc.; and the like.

Suitable example of "halo" may include fluoro, chloro, bromo and iodo.

Suitable example of "lower alkenyloxy" may include
10 vinyloxy, 1-(or 2-)propenyloxy, 1-(or 2- or 3-)butenyloxy, 1-(or 2- or 3- or 4-)pentyloxy, 1-(or 2- or 3- or 4- or 5-)hexenyloxy, and the like, in which the preferred one may be (C₂-C₆)alkenyloxy, and the most preferred one may be 5-hexenyloxy.

15 Suitable example of "higher alkenyloxy" may include (C₇-C₂₀)alkenyloxy, in which the preferred one may be 6-heptenyloxy and 7-octenyloxy.

Suitable example of "cyclo(lower)alkyl" may include cyclopropyl, cyclobutyl, cyclopentyl, cyclohexyl, and the
20 like, in which the preferred one may be cyclo(C₄-C₆)alkyl, and the most preferred one may be cyclohexyl.

Suitable example of "higher alkanoyl" may include heptanoyl, octanoyl, nonanoyl, decanoyl, undecanoyl, lauroyl, tridecanoyl, tetradecanoyl, pentadecanoyl,
25 hexadecanoyl, heptadecanoyl, octadecanoyl, nonadecanoyl, icosanoyl, and the like, in which the preferred one may be (C₇-C₂₀)alkanoyl, and the most preferred one may be hexadecanoyl.

Suitable example of "ar(lower)alkyl" may include
30 benzyl, phenethyl, phenylpropyl, phenylbutyl, phenylpentyl, phenylhexyl, naphthylmethyl, naphthylethyl, naphthylpropyl, naphthylbutyl, naphthylpentyl, naphthylhexyl, and the like, in which the preferred one may be phenyl(C₁-C₄)alkyl, and the most preferred one may
35 be benzyl.

Suitable example of "protected amino" may include lower or higher alkoxycarbonylamino (e.g., methoxycarbonylamino, ethoxycarbonylamino, t-butoxycarbonylamino, t-pentyloxycarbonylamino, heptyloxycarbonylamino, etc.), ar(lower)alkoxycarbonylamino [e.g., phenyl(lower)alkoxycarbonylamino (e.g., benzyloxycarbonylamino, etc.), etc.], an amino group substituted with a conventional protecting group such as ar(lower)alkyl which may have suitable substituent(s) (e.g., benzyl, trityl, etc.) and the like, in which the preferred one may be phenyl(lower)alkoxycarbonylamino, and the most preferred one may be benzyloxycarbonylamino.

Suitable example of "lower alkanoyl" in the term of "lower alkanoyl substituted with unsaturated 6-membered heteromonocyclic group containing at least one nitrogen atom which may have one or more suitable substituent(s)" can be referred to aforementioned "lower alkanoyl", in which the preferred one may be (C₁-C₄)alkanoyl, and the more preferred one may be formyl.

Suitable example of "unsaturated 6-membered heteromonocyclic group containing at least one nitrogen atom" in the term of "lower alkanoyl substituted with unsaturated 6-membered heteromonocyclic group containing at least one nitrogen atom which may have one or more suitable substituent(s)" may include pyridyl, dihydropyridyl, pyrimidyl, pyrazinyl, pyridazinyl, triazinyl (e.g., 4H-1,2,4-triazinyl, 1H-1,2,3-triazinyl, etc.), tetrazinyl (e.g., 1,2,4,5-tetrazinyl, 1,2,3,4-tetrazinyl, etc.), and the like, in which the preferred one may be unsaturated 6-membered heteromonocyclic group containing 1 to 3 nitrogen atom(s), and the most preferred one may be pyridyl and pyridazinyl.

Suitable example of "suitable substituent(s)" in the

term of "lower alkanoyl substituted with unsaturated 6-membered heteromonocyclic groups containing at least one nitrogen atom which may have one or more suitable substituent(s)" can be referred to aforementioned

5 "suitable substituent(s)",

in which the preferred one may be higher alkoxy, higher alkoxy(lower)alkyl, heterocyclic group which may have aryl having higher alkoxy, aryl which may have one or more higher alkoxy, aryl substituted with aryl which may have lower alkoxy, heterocyclic group which may have aryl having lower alkoxy(higher)alkoxy, and heterocyclic group which may have aryl having lower alkoxy, and the more preferred one may be (C₇-C₁₄)alkoxy, (C₇-C₁₄)alkoxy-(C₁-C₄)alkyl, 3 to 8-membered saturated heteromonocyclic group containing at least one nitrogen atom which may have phenyl having 1 to 3 (C₇-C₁₄)alkoxy, phenyl which may have 1 to 3 (C₇-C₁₄)alkoxy, phenyl substituted with phenyl which may have 1 to 3 (C₃-C₆)alkoxy, 3 to 8-membered saturated heteromonocyclic group containing at least one nitrogen atom which may have phenyl having (C₁-C₄)-alkoxy(C₇-C₁₄)alkoxy, and 3 to 8-membered saturated heteromonocyclic group containing at least one nitrogen atom which may have phenyl having 1 to 3 (C₃-C₆)alkoxy, and the most preferred one may be octyloxy, octyloxymethyl, piperazinyl which has phenyl having heptyloxy or octyloxy, phenyl having heptyloxy, phenyl substituted with phenyl having butoxy, piperazinyl which has phenyl having methoxyoctyloxy, and piperazinyl which has phenyl having hexyloxy.

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Suitable example of "lower alkanoyl" in the term of "lower alkanoyl substituted with 1,2,3,4-tetrahydroisoquinoline which may have one or more suitable substituent(s)" can be referred to aforementioned "lower alkanoyl",

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in which the preferred one may be (C₁-C₄)-
alkanoyl, and the more preferred one may be formyl.

Suitable example of "suitable substituent(s)" in the
term of "lower alkanoyl substituted with 1,2,3,4-
5 tetrahydroisoquinoline which may have one or more suitable
substituent(s)" can be referred to aforementioned
"suitable substituent(s)",

in which the preferred one may be lower alkoxy, higher
alkoxy, lower alkyl, higher alkyl and lower
10 alkoxy carbonyl, and the more preferred one may be
(C₇-C₁₄)alkoxy and (C₁-C₄)alkoxy carbonyl, and the most
preferred one may be octyloxy and tert-butoxy carbonyl.

Suitable example of "lower alkanoyl" in the term of
15 "lower alkanoyl substituted with unsaturated condensed
heterocyclic group containing at least one oxygen atom
which may have one or more suitable substituent(s)" can be
referred to aforementioned "lower alkanoyl",

in which the preferred one may be (C₁-C₄)alkanoyl, and
20 the more preferred one may be formyl.

Suitable example of "unsaturated condensed
heterocyclic group containing at least one oxygen atom" in
the term of "lower alkanoyl substituted with unsaturated
condensed heterocyclic group containing at least one
25 oxygen atom which may have one or more suitable
substituent(s)" may include unsaturated condensed
heterocyclic group containing one or more oxygen atom(s)
and, optionally, another hetero atom(s) except oxygen
atom,

in which the preferred one may be unsaturated condensed
30 heterocyclic group containing 1 to 3 oxygen atom(s),
unsaturated condensed heterocyclic group containing 1 to 2
oxygen atom(s) and 1 to 2 sulfur atom(s) and unsaturated
condensed heterocyclic group 1 to 3 oxygen atom(s) and 1
35 to 3 nitrogen atom(s), and the more preferred one may be

benzo[b]furanyl, isobenzofuranyl, chromenyl, xanthenyl, benzoxazolyl, benzoxadiazolyl, dihydrooxathiinyl, phenoxathiinyl, and the like, and the most preferred one may be benzo[b]furanyl, chromenyl and benzoxazolyl.

5 Suitable example of "suitable substituent(s)" in the term of "lower alkanoyl substituted with unsaturated condensed heterocyclic group containing at least one oxygen atom which may have one or more suitable substituent(s)" can be referred to aforementioned

10 "suitable substituent(s)",

in which the preferred one may be lower alkoxy, higher alkoxy, lower alkyl, higher alkyl, oxo, aryl which may have one or more lower alkoxy, heterocyclic group which may have one or more higher alkoxy, and aryl substituted

15 with aryl which may have one or more lower alkyl, and the more preferred one may be (C₇-C₁₄)alkoxy, (C₁-C₄)alkyl, (C₇-C₁₄)alkyl, oxo, phenyl which may have 1 to 3 (C₃-C₆)alkoxy, unsaturated 6-membered heteromonochclic group containing at least one nitrogen atom which may have

20 1 to 3 (C₇-C₁₄)alkoxy, and phenyl substituted with phenyl which may have 1 to 3 (C₃-C₆)alkyl, and the most preferred one may be octyloxy, methyl, nonyl, oxo, phenyl having hexyloxy, pyridyl having octyloxy, and phenyl substituted with phenyl having hexyl.

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Suitable example of "lower alkanoyl" in the term of "lower alkanoyl substituted with unsaturated condensed heterocyclic group containing 1 to 3 sulfur atom(s) which may have one or more suitable substituent(s)" can be

30 referred to aforementioned "lower alkanoyl",

in which the preferred one may be (C₁-C₄)alkanoyl, and the more preferred one may be formyl.

Suitable example of "unsaturated condensed heterocyclic group containing 1 to 3 sulfur atom(s)" in

35 the term of "lower alkanoyl substituted with unsaturated

condensed heterocyclic group containing 1 to 3 sulfur atom(s) which may have one or more suitable substituent(s)" may include unsaturated condensed heterocyclic group containing only 1 to 3 sulfur atom(s),
5 in which the preferred one may be benzothienyl and benzodithiynyl, and the most preferred one may be benzothienyl.

Suitable example of "suitable substituent(s)" in the term of "lower alkanoyl substituted with unsaturated condensed heterocyclic group containing 1 to 3 sulfur atom(s) which may have one or more suitable substituent(s)" can be referred to aforementioned "suitable substituent(s)",
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in which the preferred one may be lower alkoxy, higher alkoxy, lower alkyl and higher alkyl, and more preferred one may be (C₇-C₁₄)alkoxy, and the most preferred one may be octyloxy.
15

Suitable example of "lower alkanoyl" in the term of "lower alkanoyl substituted with unsaturated condensed heterocyclic group containing 2 or more nitrogen atom(s) which may have one or more suitable substituent(s)" can be referred to aforementioned "lower alkanoyl",
20

in which the preferred one may be (C₁-C₄)alkanoyl, and the most preferred one may be formyl.
25

Suitable example of "unsaturated condensed heterocyclic group containing 2 or more nitrogen atom(s)" in the term of "lower alkanoyl substituted with unsaturated condensed heterocyclic group containing 2 or more nitrogen atom(s) which may have one or more suitable substituent(s)" may include 1H-indazolyl, purinyl, phthalazinyl, benzoimidazolyl, naphthyridinyl, quinoxalinyl, quinazolyl, cinnolinyl, pteridinyl, and the like,
30

in which the most preferred one may be benzoimidazolyl.
35

Suitable example of "suitable substituent(s)" in the term of "lower alkanoyl substituted with unsaturated condensed heterocyclic group containing 2 or more nitrogen atom(s) which may have one or more suitable substituent(s)" can be referred to aforementioned

"suitable substituent(s)",

in which the preferred one may be lower alkoxy, higher alkoxy, lower alkyl, higher alkyl, aryl which may have one or more lower alkoxy and aryl which may have one or more higher alkoxy, and the more preferred one may be (C₇-C₁₄)alkyl and phenyl which may have 1 to 3 (C₁-C₆)alkoxy, and the most preferred one may be nonyl and phenyl which may have hexyloxy.

Suitable example of "lower alkanoyl" in the term of "lower alkanoyl substituted with saturated 3 to 8-membered heteromonocyclic group containing at least one nitrogen atom which may have one or more suitable substituent(s)" can be referred to aforementioned "lower alkanoyl",

in which the preferred one may be (C₁-C₄)alkanoyl, and the more preferred one may be formyl.

Suitable example of "saturated 3 to 8-membered heteromonocyclic group containing at least one nitrogen atom" in the term of "lower alkanoyl substituted with saturated 3 to 8-membered heteromonocyclic group containing at least one nitrogen atom which may have one or more suitable substituent(s)" may include pyrrolidinyl, imidazolidinyl, piperidyl, piperazinyl, pyrazolidinyl, morpholinyl, thiomorpholinyl, and the like,

in which the preferred one may be piperidyl and piperazinyl.

Suitable example of "suitable substituent(s)" in the term of "lower alkanoyl substituted with saturated 3 to 8-membered heteromonocyclic group containing at least one nitrogen atom which may have one or more suitable

substituent(s)" may include lower alkoxy, higher alkoxy,
higher alkoxy(lower)alkyl, lower alkyl, higher alkyl, oxo,
aryl which may have one or more lower alkoxy,
aryl which may have one or more higher alkoxy,
5 aryl which may have one or more lower alkyl,
aryl which may have one or more higher alkyl,
aroyl which may have one or more lower alkoxy,
aroyl which may have one or more higher alkoxy,
aroyl which may have one or more lower alkyl,
10 aroyl which may have one or more higher alkyl,
and the like,

in which the preferred one may be
aryl which may have one or more lower alkoxy,
aryl which may have one or more higher alkoxy,
15 aroyl which may have one or more lower alkoxy and
aroyl which may have one or more higher alkoxy, and the
more preferred one may be aryl which may have 1 to 3
higher alkoxy and aroyl which may have 1 to 3 higher
alkoxy, and the much more preferred one may be phenyl
20 which may have 1 to 3 (C₇-C₁₄)alkoxy and naphthoyl which
may have 1 to 3 (C₇-C₁₄)alkoxy, and the most preferred one
may be phenyl which may have octyloxy and naphthoyl which
may have heptyloxy.

25 Suitable example of "ar(lower)alkenoyl" in the term
of "ar(lower)alkenoyl substituted with aryl which may have
one or more suitable substituent(s)" may include
phenyl(lower)alkenoyl (e.g., 3-phenylacryloyl, (2- or 3-
or 4-)phenyl-(2- or 3-)butenoyl, 3-phenylmethacryloyl,
30 (2- or 3- or 4- or 5-)phenyl-(2- or 3- or 4-)pentanoyl,
(2- or 3- or 4- or 5- or 6-)phenyl-(2- or 3- or 4- or 5-)-
hexanoyl, etc.), naphthyl(lower)alkenoyl (e.g.,
3-naphthylacryloyl, (2- or 3- or 4-)naphthyl-(2- or
3-)butenoyl, (2- or 3- or 4- or 5-)naphthyl-(2- or 3- or
35 4-)pentanoyl, (2- or 3- or 4- or 5- or 6-)naphthyl-(2- or

3- or 4- or 5-)hexanoyl, etc.), and the like,
in which the preferred one may be 3-phenylacryloyl and 3-methyl-3-phenylacryloyl.

5 Suitable example of "suitable substituent(s)" in the term of "ar(lower)alkenoyl substituted with aryl which may have one or more suitable substituent(s)" can be referred to aforementioned "suitable substituent(s)",

in which the preferred one may be lower alkoxy, lower alkyl, higher alkyl, lower alkoxy(lower)alkyl,
10 halo(lower)alkoxy, lower alkenyloxy, halo(higher)alkoxy, and lower alkoxy(higher)alkoxy and the much more preferred one may be (C₁-C₆)alkoxy, (C₁-C₆)alkyl, (C₇-C₁₄)alkyl, (C₁-C₄)alkoxy(C₃-C₆)alkyl, halo(C₃-C₆)alkoxy, (C₃-C₆)alkenyloxy, halo(C₇-C₁₄)alkoxy, and (C₁-C₄)alkoxy(C₇-
15 C₁₄)alkoxy and the most preferred one may be pentyloxy, heptyl, pentyl, methoxyhexyl, fluorohexyloxy, isohexyloxy, 5-hexenyloxy, haloheptyloxy, methoxyheptyloxy, methoxyoctyloxy, and butyloxy.

20 Suitable example of "naphthyl(lower)alkenoyl" in the term of "naphthyl(lower)alkenoyl which may have one or more higher alkoxy" may include 3-naphthylacryloyl, (2- or 3- or 4-)naphthyl-(2- or 3-)butenoyl, (2- or 3- or 4- or 5-)naphthyl-(2- or 3- or 4-)pentanoyl, (2- or 3- or 4- or 5- or 6-)naphthyl-(2- or 3- or 4- or 5-)hexanoyl, and the
25 like,
in which the preferred one may be 3-naphthylacryloyl.

30 Suitable example of "lower alkynoyl" in the term of "lower alkynoyl which may have one or more suitable substituent(s)" may include 2-propynoyl, (2- or 3-)butynoyl, (2- or 3- or 4-)pentynoyl, (2- or 3- or 4- or 5-)hexynoyl, and the like,
in which the preferred one may be 2-propynoyl.

35 Suitable example of "suitable substituent(s)" in the

term of "lower alkynoyl which may have one or more suitable substituent(s)" can be referred to aforementioned "suitable substituent(s)",

in which the preferred one may be aryl which may have one or more lower alkoxy, aryl which may have one or more higher alkoxy, aryl substituted with aryl which may have one or more lower alkyl and aryl substituted with aryl which may have one or more higher alkyl, and the more preferred one may be aryl substituted with aryl which may have 1 to 3 lower alkyl and aryl which may have 1 to 3 higher alkoxy, and the much more preferred one may be phenyl substituted with phenyl which may have 1 to 3 (C₁-C₆)alkyl and phenyl which may have 1 to 3 (C₇-C₁₄)alkoxy, and the most preferred one may be phenyl substituted with phenyl which may have pentyl and naphthyl which may have heptyloxy.

Suitable example of "ar(C₂-C₆)alkanoyl" in the term of "ar(C₂-C₆)alkanoyl substituted with aryl having one or more suitable substituent(s), in which ar(C₂-C₆)alkanoyl may have one or more suitable substituent(s)" may include phenyl(C₂-C₆)alkanoyl [e.g., phenylacetyl, (2- or 3-)-phenylpropanoyl, (2- or 3- or 4-)-phenylbutanoyl, (2- or 3- or 4- or 5-)-phenylpentanoyl, (2- or 3- or 4- or 5- or 6-)-phenylhexanoyl, etc.], naphthyl(C₂-C₆)alkanoyl [e.g. naphthylacetyl, (2- or 3-)-naphthylpropanoyl, (2- or 3- or 4-)-naphthylbutanoyl, (2- or 3- or 4- or 5-)-naphthylpentanoyl, (2- or 3- or 4- or 5- or 6-)-naphthylhexanoyl, etc.], and the like, in which the preferred one may be 2-phenylacetyl and 3-phenylpropanoyl.

Suitable example of "suitable substituent(s)" in the term of "ar(C₂-C₆)alkanoyl substituted with aryl having one or more suitable substituent(s), in which ar(C₂-C₆)-alkanoyl may have one or more suitable substituent(s)" may

include lower alkoxy, higher alkoxy, lower alkyl, higher alkyl, higher alkoxy(lower)alkyl, oxo, aryl having one or more lower alkoxy, aryl having one or more higher alkoxy, aryl having one or more lower alkyl, aryl having one or more higher alkyl, aryl substituted with aryl having one or more lower alkoxy, aryl substituted with aryl having one or more higher alkoxy, aryl substituted with aryl having one or more lower alkyl, aryl substituted with aryl having one or more higher alkyl, aryl having one or more lower alkoxy(lower)alkoxy and the like,

in which the preferred one may be lower alkoxy, higher alkoxy, lower alkyl, higher alkyl, and phenyl having 1 to 3 lower alkoxy(lower)alkoxy and the much more preferred one may be (C₁-C₆)alkoxy, (C₁-C₆)alkyl, (C₇-C₁₄)alkyl and phenyl having (C₁-C₄)alkoxy(C₃-C₆)alkoxy and the most preferred one may be pentyloxy, pentyl, heptyl and phenyl having methoxypentyloxy.

Suitable example of "suitable substituent(s)" in the term of "in which ar(C₂-C₆)alkanoyl may have one or more suitable substituent(s)" may be hydroxy, oxo, amino and aforementioned "protected amino".

Suitable example of "(C₂-C₆)alkanoyl" in the term of "(C₂-C₆)alkanoyl substituted with naphthyl having higher alkoxy" may include acetyl, propanoyl, butanoyl, pentanoyl, hexanoyl, and the like,

in which the preferred one may be propanoyl.

Suitable example of "higher alkoxy" in the term of "(C₂-C₆)alkanoyl substituted with naphthyl having higher alkoxy" can be referred to aforementioned "higher alkoxy", in which the preferred one may be (C₇-C₁₄)alkoxy, and the most preferred one may be heptyloxy.

Suitable example of "aroyl" in the term of "aroyl substituted with heterocyclic group which may have one or

more suitable substituent(s), in which aroyl may have one or more suitable substituent(s)" may include benzoyl, toluoyl, naphthoyl, and the like,

in which the preferred one may be benzoyl.

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Suitable example of "heterocyclic group" in the term of "aroyl substituted with heterocyclic group which may have one or more suitable substituent(s), in which aroyl may have one or more suitable substituent(s)" may include
10 unsaturated 3 to 8-membered (more preferably 5 or 6-membered) heteromonocyclic group containing 1 to 4 nitrogen atom(s), for example, pyrrolyl, pyrrolinyl, imidazolyl, pyrazolyl, pyridyl, dihydropyridyl, pyrimidyl, pyrazinyl, pyridazinyl, triazolyl (e.g., 4H-1,2,4-
15 triazolyl, 1H-1,2,3-triazolyl, 2H-1,2,3-triazolyl, etc.), tetrazolyl (e.g., 1H-tetrazolyl, 2H-tetrazolyl, etc.), etc.;

saturated 3 to 8-membered (more preferably 5 or 6-membered) heteromonocyclic group containing 1 to 4
20 nitrogen atom(s), for example, pyrrolidinyl, imidazolidinyl, piperidyl, piperazinyl, etc.;

unsaturated condensed heterocyclic group containing 1 to 4 nitrogen atom(s), for example, indolyl, isoindolyl, indolinyl, indolizinyll, benzimidazolyl, quinolyl,
25 isoquinolyl, indazolyl, benzotriazolyl, etc.;

unsaturated 3 to 8-membered (more preferably 5 or 6-membered) heteromonocyclic group containing 1 to 2 oxygen atom(s) and 1 to 3 nitrogen atom(s), for example, oxazolyl, isoxazolyl, oxadiazolyl (e.g.,
30 1,2,4-oxadiazolyl, 1,3,4-oxadiazolyl, 1,2,5-oxadiazolyl, etc.), etc.;

saturated 3 to 8-membered (more preferably 5 or 6-membered) heteromonocyclic group containing 1 to 2 oxygen atom(s) and 1 to 3 nitrogen atom(s), for example,
35 morpholinyl, sydnonyl, etc.;

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an oxygen atom and 1 to 2 sulfur atom(s), for example, benzoxathiinyl, etc.; and the like,

in which the preferred one may be saturated 3 to 8-membered heteromonocyclic group containing 1 to 4 nitrogen atom(s), unsaturated 3 to 8-membered heteromonocyclic group containing 1 to 2 oxygen atom(s) and 1 to 3 nitrogen atom(s),

unsaturated 3 to 8-membered heteromonocyclic group containing 1 to 4 nitrogen atom(s), and

unsaturated 3 to 8-membered heteromonocyclic group containing 1 to 2 sulfur atom(s) and 1 to 3 nitrogen atom(s), and the most preferred one may be piperazinyl, isoxazolyl, oxadiazolyl, thiadiazolyl, pyrazolyl, piperidyl, oxazolyl and pyrimidyl.

Suitable example of "suitable substituent(s)" in the term of "aroyl substituted with heterocyclic group which may have one or more suitable substituent(s), in which aroyl may have one or more suitable substituent(s)", can be referred to aforementioned "suitable substituent(s)",

in which the preferred one may be aryl which may have 1 to 3 higher alkoxy, aryl which may have 1 to 3 lower alkoxy, higher alkyl, heterocyclic group, aryl which may have 1 to 3 lower alkoxy(higher)alkoxy, aryl which may have higher alkenyloxy, heterocyclic group which may have aryl having lower alkoxy, cyclo(lower)alkyl which may have aryl, aryl which may have 1 to 3 lower alkyl, aryl which may have cyclo(lower)alkyl, aryl which may have higher alkenyloxy, aryl substituted with heterocyclic group which may have lower alkyl and oxo, cyclo(lower)alkyl which may have lower alkyl, aryl substituted with aryl which may have 1 to 3 lower alkoxy, and aryl which may have heterocyclic group, and the more preferred one may be phenyl which may have 1 to 3 (C₇-C₁₄)alkoxy, phenyl which may have 1 to 3 (C₃-C₆)alkoxy, (C₇-C₁₄)alkyl, saturated 3

to 8-membered heteromonocyclic group containing 1 to 4
nitrogen atom(s), phenyl which may have 1 to 3 (C₁-
C₄)alkoxy (C₇-C₁₄)alkoxy, phenyl which may have (C₇-
C₁₄)alkenyloxy, saturated 3 to 8-membered heteromonocyclic
5 group containing 1 to 4 nitrogen atom(s) substituted with
phenyl having (C₃-C₆)alkoxy, cyclo(C₃-C₆)alkyl which may
have phenyl, phenyl which may have 1 to 3 (C₃-C₆)alkyl,
phenyl which may have cyclo(C₃-C₆)alkyl, phenyl which may
have (C₇-C₁₄)alkenyloxy, phenyl substituted with
10 heterocyclic group which may have (C₃-C₆)alkyl and oxo,
cyclo(C₃-C₆)alkyl which may have (C₃-C₆)alkyl, phenyl
substituted with phenyl which may have 1 to 3 (C₁-
C₄)alkoxy, and phenyl which may have 3 to 8-membered
heteromonocyclic group containing 1 to 4 nitrogen atom(s),
15 and the most preferred one may be phenyl having octyloxy,
phenyl having pentyloxy, phenyl having hexyloxy, heptyl,
piperidyl, phenyl having isohexyloxy, phenyl having
heptyloxy, phenyl having methoxyheptyloxy, phenyl having
methoxyoctyloxy, phenyl having 6-heptenyloxy, piperidyl
20 substituted with phenyl having hexyloxy, cyclohexyl having
phenyl, phenyl having hexyl, phenyl having cyclohexyl,
phenyl having 7-octenyloxy, phenyl substituted with
triazolyl having lower alkyl and oxo, cyclohexyl having
pentyl, phenyl having methoxyoctyloxy, nonyl, phenyl
25 substituted with phenyl having propoxy, and phenyl having
piperidine.

Suitable example of "suitable substituent(s)" in the
term of "in which aroyl may have one or more suitable
substituent(s)" may be halogen, in which the preferred one
30 may be fluoro and chloro.

Suitable example of "aroyl" in the term of "aroyl
substituted with aryl having heterocyclic(higher)alkoxy,
in which heterocyclic group may have one or more suitable
35 substituent(s)" may include benzoyl, toluoyl, naphthoyl,

anthrylcarbonyl and the like,

in which the preferred one may be benzoyl.

Suitable example of "heterocyclic" moiety in the term of "aroyl substituted with aryl having

5 heterocyclic(higher)alkoxy, in which heterocyclic group may have one or more suitable substituent(s)" can be referred to the ones as exemplified before for

"heterocyclic group" in the term of "aroyl substituted with heterocyclic group which may have one or more
10 suitable substituent(s)",

in which the preferred one may be unsaturated 3 to 8-membered heteromonocyclic group containing 1 to 4 nitrogen atom(s) and saturated 3 to 8-membered heteromonocyclic group containing 1 to 2 oxygen atom(s) and 1 to 3 nitrogen
15 atom(s), and the most preferred one may be triazolyl, tetrazolyl and morpholinyl.

Suitable example of "(higher)alkoxy" moiety in the term of "aroyl substituted with aryl having heterocyclic(higher)alkoxy, in which heterocyclic group
20 may have one or more suitable substituent(s)" can be referred to aforementioned "higher alkoxy",

in which the preferred one may be (C₇-C₁₄)alkoxy, and the most preferred one may be octyloxy.

Suitable example of "aryl" in the term of "aroyl substituted with aryl having heterocyclic(higher)alkoxy, in which heterocyclic group may have one or more suitable substituent(s)" can be referred to aforementioned "aryl",
25 in which the preferred one may be phenyl.

Suitable example of "suitable substituent(s)" in the term of "in which heterocyclic group may have one or more suitable substituent(s)" may be lower alkyl, in which the
30 preferred one may be methyl.

Suitable example of "aroyl" in the term of "aroyl substituted with aryl having lower alkoxy(higher)alkoxy"
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in which the preferred one may be benzoyl.

Suitable example of "lower alkoxy(higher)alkoxy" in the term of "aryl substituted with aryl having lower alkoxy(higher)alkoxy" may be methoxyheptyloxy,

20 Suitable example of "aroyl" in the term of "aroyl substituted with aryl having lower alkenyl(lower)alkoxy" may include benzoyl, toluoyl, naphthoyl, anthrylcarbonyl and the like.

in which the preferred one may be benzoyl.

25 Suitable example of "aryl" in the term of "aroyl
substituted with aryl having lower alkenyl(lower)alkoxy"
can be referred to aforementioned "aryl",
in which the preferred one may be phenyl.

Suitable example of "lower alkenyl(lower)alkoxy" in the term of "aroyl substituted with aryl having lower alkenyl(lower)alkoxy" may be vinylmethoxy, vinyllethoxy, vinylpropoxy, vinylbutoxy, vinylpentyloxy, vinylhexyloxy, 1-(or 2-)propenylmethoxy, 1-(or 2-)propenylethoxy, 1-(or 2-)propenylpropoxy, 1-(or 2-)propenylbutoxy, 1-(or 2-)propenylpentyloxy, 1-(or 2-)propenylhexyloxy, 1-(or 2- or

3-)butenylbutoxy, 1-(or 2- or 3-)butenylhexyloxy, 1-(or 2-
or 3- or 4-)pentenylpentyloxy, 1-(or 2- or 3- or 4-)-
pentenylhexyloxy, 1-(or 2- or 3- or 4- or 5-)-
hexenylbutoxy, 1-(or 2- or 3- or 4- or 5-)hexenylhexyloxy,
5 or the like,

in which the preferred one may be (C₂-C₆)alkenyl(C₁-
C₆)alkoxy, and the more preferred one may be
vinylhexyloxy.

10 Suitable example of "aroyl substituted with 2 lower
alkoxy" may include benzoyl substituted with 2 lower
alkoxy and naphthoyl substituted with 2 lower alkoxy,
in which the preferred one may be benzoyl substituted
with 2 (C₁-C₆)alkoxy, and the most preferred one may be
15 benzoyl substituted with 2 pentyloxy.

Suitable example of "aroyl substituted with aryl
having lower alkyl" may include benzoyl substituted with
phenyl having lower alkyl, benzoyl substituted with
20 naphthyl having lower alkyl, naphthoyl substituted with
phenyl having lower alkyl, naphthoyl substituted with
naphthyl having lower alkyl, and the like,
in which the preferred one may be benzoyl substituted
with phenyl having (C₁-C₆)alkyl, and the most preferred
25 one may be benzoyl substituted with phenyl having hexyl
and benzoyl substituted with phenyl having pentyl.

Suitable example of "aroyl substituted with aryl
having higher alkyl" may include benzoyl substituted with
30 phenyl having higher alkyl, benzoyl substituted with
naphthyl having higher alkyl, naphthoyl substituted with
phenyl having higher alkyl, naphthoyl substituted with
naphthyl having higher alkyl, and the like,
in which the preferred one may be benzoyl substituted
35 with phenyl having (C₇-C₁₄)alkyl, and the most preferred

one may be benzoyl substituted with phenyl having heptyl.

Suitable example of "aryloxy" moiety in the term of "aryloxy(lower)alkanoyl which may have one or more suitable substituent(s)" may include phenoxy, mesityloxy, tolyloxy, naphthyloxy, anthryloxy, and the like, in which the preferred one may be phenoxy.

Suitable example of "lower alkanoyl" moiety in the term of "aryloxy(lower)alkanoyl which may have one or more suitable substituent(s)" can be referred to aforementioned "lower alkanoyl",

in which the preferred one may be formyl, acetyl, 2,2-dimethylacetyl, propionyl, butyryl, isobutyryl and pentanoyl, hexanoyl, and the more preferred one may be (C₁-C₆)alkanoyl, and the much more preferred one may be formyl, acetyl, propionyl and 2,2-dimethylacetyl.

Suitable example of "suitable substituent(s)" in the term of "aryloxy(lower)alkanoyl which may have one or more suitable substituent(s)" can be referred to aforementioned "suitable substituent(s)",

in which the preferred one may be (C₇-C₁₄)alkoxy, and the more preferred one may be octyloxy.

Suitable example of "ar(lower)alkoxy" moiety in the term of "ar(lower)alkoxy(lower)alkanoyl which may have one or more suitable substituent(s)" may include phenyl(lower)alkoxy [e.g., phenylmethoxy, (1- or 2-)-phenylethoxy, phenylpropoxy, 2-phenyl-1-methylpropoxy, 3-phenyl-2,2-dimethylpropoxy, (1- or 2- or 3- or 4-)-phenylbutoxy, (1- or 2- or 3- or 4- or 5-)-phenylpentyloxy, (1- or 2- or 3- or 4- or 5- or 6-)-phenylhexyloxy, etc.], naphthyl(lower)alkoxy [e.g. naphthylmethoxy, (1- or 2-)-naphthylethoxy, 1-naphthylpropoxy, 2-naphthyl-1-methylpropoxy, 3-naphthyl-2,2-dimethylpropoxy, (1- or 2- or 3- or 4-)-naphthylbutoxy,

(1- or 2- or 3- or 4- or 5-)naphthylpentyloxy, (1- or 2- or 3- or 4- or 5- or 6-)naphthylhexyloxy, etc.], and the like,

5 in which the preferred one may be naphthyl(C₁-C₄)alkoxy, and the more preferred one may be naphthylmethoxy.

Suitable example of "(lower)alkanoyl" moiety in the term of "ar(lower)alkoxy(lower)alkanoyl which may have one or more suitable substituent(s)" can be referred to aforementioned "lower alkanoyl",

10 in which the preferred one may be (C₁-C₄)alkanoyl, and the more preferred one may be formyl.

Suitable example of "suitable substituent(s)" in the term of "ar(lower)alkoxy(lower)alkanoyl which may have one or more suitable substituent(s)" can be referred to aforementioned "suitable substituent(s)",

15 in which the preferred one may be lower alkoxy, higher alkoxy, lower alkyl and higher alkyl, and the more preferred one may be higher alkoxy, and the much more preferred one may be (C₇-C₁₄)alkoxy, and the most
20 preferred one may be heptyloxy.

Suitable example of "arylamino" moiety in the term of "arylamino(lower)alkanoyl which may have one or more suitable substituent(s)" may include phenylamino, mesitylamino, tolylamino, naphthylamino, anthrylamino and the like,

25 in which the preferred one may be phenylamino and naphthylamino.

Suitable example of "lower alkanoyl" moiety in the term of "arylamino(lower)alkanoyl which may have one or more suitable substituent(s)" can be referred to aforementioned "lower alkanoyl",

30 in which the preferred one may be (C₁-C₄)alkanoyl, and the more preferred one may be formyl.

35 Suitable example of "suitable substituent(s)" in the

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in which the preferred one may be lower alkoxy, higher alkoxy, lower alkyl, higher alkyl, aryl which may have 1 to 3 lower alkoxy and aryl which may have 1 to 3 higher alkoxy, and the more preferred one may be (C₇-C₁₄)alkoxy, and phenyl which may have 1 to 3 (C₇-C₁₄)alkoxy, and the most preferred one may be heptyloxy and phenyl which may have heptyloxy.

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the term of "lower alkoxy(higher)alkanoyl, in which higher alkanoyl may have one or more suitable substituent(s)" may be (C₁-C₄)alkoxy(C₇-C₂₀)alkanoyl, in which the preferred one may be methoxyoctadecanoyl.

5 Suitable example of "suitable substituent(s)" in the term of "lower alkoxy(higher)alkanoyl, in which higher alkanoyl may have one or more suitable substituent(s)" may be amino and aforementioned "protected amino", in which the preferred one may be amino and
10 ar(lower)alkoxycarbonylamino, and the most preferred one may be amino and benzyloxycarbonylamino.

15 Suitable example of "aroyl" in the term of "aroyl substituted with aryl having heterocyclicoxy, in which heterocyclicoxy may have one or more suitable substituent(s)" can be referred to aforementioned "aroyl", in which the preferred one may be benzoyl.

20 Suitable example of "aryl" in the term of "aroyl substituted with aryl having heterocyclicoxy, in which heterocyclicoxy may have one or more suitable substituent(s)" can be referred to aforementioned "aryl", in which the preferred one may be phenyl.

25 Suitable example of "heterocyclic" moiety in the term of "aroyl substituted with aryl having heterocyclicoxy, in which heterocyclicoxy may have one or more suitable substituent(s)" can be referred to aforementioned "heterocyclic" moiety, in which the preferred one may be unsaturated 3 to 8-membered heteromonocyclic group containing 1 to 4 nitrogen atom(s), and the most preferred
30 one may be pyridazinyl.

35 Suitable example of "suitable substituent(s)" in the term of "aroyl substituted with aryl having heterocyclicoxy, in which heterocyclicoxy may have one or more suitable substituent(s)" may be aryl, in which the preferred one may be phenyl.

substituted with aryl having lower
alkoxy(lower)alkoxy(higher)alkoxy", "aroyle substituted
with aryl having lower alkoxy(lower)alkoxy", "aroyle
substituted with aryl which has aryl having lower alkoxy",
5 "aroyle substituted with aryl which has aryl having lower
alkoxy(lower)alkoxy", "aroyle substituted with aryl having
heterocyclicoxy(higher)alkoxy", "aroyle substituted with
aryl having aryloxy(lower)alkoxy" and "aroyle substituted
with aryl having heterocycliccarbonyl(higher)alkoxy" can
10 be referred to aforementioned "aroyle", in which the
preferred one may be benzoyl.

Suitable example of "aryl" in abovementioned terms
can be referred to aforementioned "aryl", in which the
preferred one may be phenyl.

15 Suitable example of "lower alkoxy(lower)alkoxy-
(higher)alkoxy" in the term of "aroyle substituted with
aryl having lower alkoxy(lower)alkoxy(higher)alkoxy" may
be (C₁-C₄)alkoxy(C₁-C₄)alkoxy(C₇-C₁₄)alkoxy, in which the
20 preferred one may be ethoxyethoxyoctyloxy.

Suitable example of "lower alkoxy(lower)alkoxy" in
the term of "aroyle substituted with aryl having lower
alkoxy(lower)alkoxy" may be (C₁-C₄)alkoxy(C₃-C₆)alkoxy, in
25 which the preferred one may be propoxyhexyloxy.

Suitable example of "lower alkoxy" in the term of
"aroyle substituted with aryl which has phenyl having lower
alkoxy" may be (C₃-C₆)alkoxy, in which the preferred one
30 may be butoxy.

Suitable example of "lower alkoxy(lower)alkoxy" in
the term of "aroyle substituted with aryl which has phenyl
having lower alkoxy(lower)alkoxy" may be (C₁-C₄)alkoxy-
35 (C₃-C₆)alkoxy, in which the preferred one may be

methoxypentyloxy and methoxyhexyloxy.

Suitable example of "heterocyclic" moiety in the term of "aroyl substituted with aryl having heterocyclicoxy(higher)alkoxy" can be referred to aforementioned "heterocyclic" moiety, in which the preferred one may be saturated 3 to 8-membered heteromonocyclic group containing an oxygen atom, and the most preferred one may be tetrahydropyranyl.

Suitable example of "higher alkoxy" moiety in the term of "aroyl substituted with aryl having heterocyclicoxy(higher)alkoxy" may be (C₇-C₁₄)alkoxy, in which the preferred one may be octyloxy.

Suitable example of "aryloxy(lower)alkoxy" in the term of "aroyl substituted with aryl having aryloxy(lower)alkoxy" may be phenoxy(C₃-C₆)alkoxy, in which the preferred one may be phoxypentyloxy.

Suitable example of "heterocyclic" moiety in the term of "aroyl substituted with aryl having heterocycliccarbonyl(higher)alkoxy" can be referred to aforementioned "heterocyclic" moiety, in which the preferred one may be saturated 3 to 8-membered heteromonocyclic group containing 1 to 4 nitrogen atom(s), and the most preferred one may be piperidyl.

Suitable example of "higher alkoxy" moiety in the term of "aroyl substituted with aryl having heterocycliccarbonyl(higher)alkoxy" can be referred to aforementioned "higher alkoxy", in which the preferred one may be (C₇-C₁₄)alkoxy, and the most preferred one may be heptyloxy.

Suitable example of "lower alkanoyl" in the term of "lower alkanoyl substituted with oxazolyl which has aryl

Suitable example of "aryl" in the term of "lower
5 alkanoyl substituted with oxazolyl which has aryl having
higher alkoxy" can be referred to aforementioned "aryl",
in which the preferred one may be phenyl.

15 Suitable example of "lower alkanoyl" in the term of
"lower alkanoyl substituted with furyl which has aryl
substituted with aryl having lower alkoxy" can be referred
to aforementioned "lower alkanoyl", in which the preferred
one may be (C₁-C₄)alkanoyl, and the most preferred one may
20 be formyl.

Suitable example of "lower alkoxy" in the term of "lower alkanoyl substituted with furyl which has aryl substituted with aryl having lower alkoxy" can be referred to aforementioned "lower alkoxy", in which the preferred one may be (C₁-C₄)alkoxy, and the most preferred one may be butoxy.

Suitable example of "lower alkanoyl" in the term of "lower alkanoyl substituted with triazolyl which has oxo and aryl having higher alkyl" can be referred to

pentanoyl, hexanoyl, and the like, in which the preferred one may be acetyl and propanoyl.

Suitable example of "aryl" in the term of "(C₂-C₆)alkanoyl substituted with aryl having higher alkoxy, in which (C₂-C₆)alkanoyl may have amino or protected amino" can be referred to aforementioned "aryl", in which the preferred one may be phenyl.

Suitable example of "higher alkoxy" in the term of "(C₂-C₆)alkanoyl substituted with aryl having higher alkoxy, in which (C₂-C₆)alkanoyl may have amino or protected amino" can be referred to aforementioned "higher alkoxy", in which the preferred one may be (C₇-C₁₄)alkoxy, and the most preferred one may be octyloxy.

Suitable example of "protected amino" in the term of "(C₂-C₆)alkanoyl substituted with aryl having higher alkoxy, in which (C₂-C₆)alkanoyl may have amino or protected amino" can be referred to aforementioned "protected amino", in which the preferred one may be ar(lower)alkoxycarbonylamino, and the most preferred one may be benzyloxycarbonylamino.

The process for preparing the object polypeptide compound [I] or a salt thereof of the present invention are explained in detail in the following.

Process 1

The object polypeptide compound [I] or a salt thereof can be prepared by reacting the compound [II] or its reactive derivative at the amino group or a salt thereof with the compound [III] or its reactive derivative at the carboxy group or a salt thereof.

Suitable reactive derivative at the carboxy group of the compound [III] may include an acid halide, an acid anhydride, an activated amide, an activated ester, and the like. Suitable examples of the reactive derivatives may

be an acid chloride; an acid azide; a mixed acid anhydride with an acid such as substituted phosphoric acid [e.g., dialkylphosphoric acid, phenylphosphoric acid, diphenylphosphoric acid, dibenzylphosphoric acid, halogenated phosphoric acid, etc.], dialkylphosphorous acid, sulfurous acid, thiosulfuric acid, sulfuric acid, sulfonic acid [e.g., methanesulfonic acid, etc.], aliphatic carboxylic acid [e.g., acetic acid, propionic acid, butyric acid, isobutyric acid, pivalic acid, pentanoic acid, isopentanoic acid, 2-ethylbutyric acid, trichloroacetic acid, etc.]; or aromatic carboxylic acid [e.g., benzoic acid, etc.]; a symmetrical acid anhydride; an activated amide with imidazole, 4-substituted imidazole, dimethylpyrazole, triazole, tetrazole or 1-hydroxy-1H-benzotriazole; or an activated ester [e.g., cyanomethyl ester, methoxymethyl ester, dimethyliminomethyl $[(CH_3)_2N^+=CH-]$ ester, vinyl ester, propargyl ester, p-nitrophenyl ester, 2,4-dinitrophenyl ester, trichlorophenyl ester, pentachlorophenyl ester, mesylphenyl ester, phenylazophenyl ester, phenyl thioester, p-nitrophenyl thioester, p-cresyl thioester, carboxymethyl thioester, pyranyl ester, pyridyl ester, piperidyl ester, 8-quinolyl thioester, etc.], or an ester with a N-hydroxy compound [e.g. N,N-dimethylhydroxylamine, 1-hydroxy-2-(1H)-pyridone, N-hydroxysuccinimide, N-hydroxyphthalimide, 1-hydroxy-1H-benzotriazole, etc.], and the like. These reactive derivatives can optionally be selected from them according to the mind of the compound [III] to be used.

Suitable salts of the compound [III] and its reactive derivative can be referred to the ones as exemplified for the object polypeptide compound [I].

The reaction is usually carried out in a conventional solvent such as water, alcohol [e.g., methanol, ethanol,

etc.], acetone, dioxane, acetonitrile, chloroform, methylene chloride, ethylene chloride, tetrahydrofuran, ethyl acetate, N,N-dimethylformamide, pyridine or any other organic solvent which does not adversely influence the reaction. These conventional solvent may also be used in a mixture with water.

In this reaction, when the compound [III] is used in a free acid form or its salt form, the reaction is preferably carried out in the presence of a conventional condensing agent such as N,N'-dicyclohexylcarbodiimide; N-cyclohexyl-N'-morpholinoethylcarbodiimide; N-cyclohexyl-N'-(4-diethylaminocyclohexyl)carbodiimide; N,N'-diethylcarbodiimide, N,N'-diisopropylcarbodiimide; N-ethyl-N'-(3-dimethylaminopropyl)carbodiimide, N,N-carbonylbis-(2-methylimidazole); pentamethyleneketene-N-cyclohexylimine; diphenylketene-N-cyclohexylimine; ethoxyacetylene; 1-alkoxy-2-chloroethylene; trialkyl phosphite; ethyl polyphosphate; isopropyl polyphosphate; phosphorus oxychloride (phosphoryl chloride); phosphorus trichloride; thionyl chloride; oxalyl chloride; lower alkyl haloformate [e.g., ethyl chloroformate, isopropyl chloroformate, etc.]; triphenylphosphine; 2-ethyl-7-hydroxybenzisoxazolium salt; 2-ethyl-5-(m-sulfophenyl)isoxazolium hydroxide intramolecular salt; 1-(p-chlorobenzenesulfonyloxy)-6-chloro-1H-benzotriazole; so-called Vilsmeier reagent prepared by the reaction of N,N-dimethylformamide with thionyl chloride, phosgene, trichloromethyl chloroformate, phosphorous oxychloride, methanesulfonyl chloride, etc.; or the like.

The reaction may also be carried out in the presence of an inorganic or organic base such as an alkali metal carbonate, alkali metal bicarbonate, tri(lower)alkylamine, pyridine, di(lower)alkylaminopyridine (e.g.,

4-dimethylaminopyridine, etc.), N-(lower)alkylmorpholine, N,N-di(lower)alkylbenzylamine, or the like.

The reaction temperature is not critical, and the reaction is usually carried out under cooling to warming.

5

The starting compound [II] is a known compound. It can be prepared by fermentation and synthetic processes disclosed in EP 0462531 A2.

10 A culture of Coleophoma sp. F-11899, which is used in said fermentation process, has been deposited with National Institute of Bioscience and Human-Technology Agency of Industrial Science and Technology (former name: Fermentation Research Institute Agency of Industrial Science and Technology) (1-3, Higashi 1-chome, Tsukuba-shi, IBARAKI 305, JAPAN) on October 26, 1989 under the
15 number of FERM BP-2635.

20 The compounds obtained by the above Process 1 can be isolated and purified by a conventional method such as pulverization, recrystallization, column-chromatography, high-performance liquid chromatography (HPLC), reprecipitation, or the like.

25 The compounds obtained by the above Process 1 may be obtained as its hydrate, and its hydrate is included within the scope of this invention.

30 It is to be noted that each of the object compound (I) may include one or more stereoisomer such as optical isomer(s) and geometrical isomer(s) due to asymmetric carbon atom(s) and double bond(s) and all such isomers and mixture thereof are included within the scope of this invention.

35

Biological property of the polypeptide
compound [I] of the present invention.

5 In order to show the usefulness of the polypeptide
compound [I] of the present invention, the biological data
of the representative compound is explained in the
following.

Test 1 (Antimicrobial activity) :

10 In vitro antimicrobial activity of the compound of
Example 17 disclosed later was determined by the two-fold
agar-plate dilution method as described below.

Test Method

15 One loopful of an overnight culture of each test
microorganism in Sabouraud broth containing 2% Glucose
(10^5 viable cells per ml) was streaked on yeast nitrogen
base dextrose agar (YNBDA) containing graded
concentrations of the object polypeptide compound [I], and
20 the minimal inhibitory concentration (MIC) was expressed
in terms of $\mu\text{g/ml}$ after incubation at 30°C for 24 hours.

Test Result

MIC ($\mu\text{g/ml}$)

25	Test compound Test organism	The compound of <u>Example 17</u>
	candida albicans FP-633	0.2

30 From the test result, it is realized that the object
polypeptide compound [I] of the present invention has an
antimicrobial activity (especially, antifungal activity).

35 The pharmaceutical composition of the present

invention can be used in the form of a pharmaceutical preparation, for example, in solid, semisolid or liquid form, which contains the object polypeptide compound (I) or a pharmaceutically acceptable salt thereof, as an
5 active ingredient in admixture with an organic or inorganic carrier or excipient which is suitable for rectal; pulmonary (nasal or buccal inhalation); ocular; external (topical); oral administration; parenteral (including subcutaneous, intravenous and intramuscular)
10 administrations; insufflation (including aerosols from metered dose inhalator); nebulizer; or dry powder inhalator.

The active ingredient may be compounded, for example,
15 with the usual non-toxic, pharmaceutically acceptable carriers in a solid form such as granules, tablets, dragees, pellets, troches, capsules, or suppositories; creams, ointments; aerosols; powders for insufflation; in a liquid form such as solutions, emulsions, or
20 suspensions for injection; ingestion; eye drops; and any other form suitable for use. And, if necessary, there may be included in the above preparation auxiliary substance such as stabilizing, thickening, wetting, emulsifying and coloring agents; perfumes or buffer; or any other commonly
25 may be used as additives.

The object polypeptide compound [I] or a pharmaceutically acceptable salt thereof is/are included in the pharmaceutical composition in an amount sufficient to produce the desired antimicrobial effect upon the
30 process or condition of diseases.

For applying the composition to human, it is preferable to apply it by intravenous, intramuscular, pulmonary, oral administration, or insufflation. While the dosage of therapeutically effective amount of the
35 object polypeptide compound [I] varies from and also

depends upon the age and condition of each individual patient to be treated, in the case of intravenous administration, a daily dose of 0.01-20 mg of the object polypeptide compound [I] per kg weight of human being in the case of intramuscular administration, a daily dose of 0.1-20 mg of the object polypeptide compound [I] per kg weight of human being, in case of oral administration, a daily dose of 0.5-50 mg of the object polypeptide compound [I] per kg weight of human being is generally given for treating or preventing infectious diseases.

Especially in case of the treatment of prevention of Pneumocystis carinii infection, the followings are to be noted.

For administration by inhalation, the compounds of the present invention are conveniently delivered in the form of an aerosol spray presentation from pressurized as powders which may be formulated and the powder compositions may be inhaled with the aid of an insufflation powder inhaler device. The preferred delivery system for inhalation is a metered dose inhalation aerosol, which may be formulated as a suspension or solution of compound in suitable propellants such as fluorocarbons or hydrocarbons.

Because of desirability to directly treat lung and bronchi, aerosol administration is a preferred method of administration. Insufflation is also a desirable method, especially where infection may have spread to ears and other body cavities.

Alternatively, parenteral administration may be employed using drip intravenous administration.

The following Preparations and Examples are given for the purpose of illustrating the present invention in more detail.

Preparation 1

To a suspension of 1-(4-Hydroxyphenyl)-4-tert-butoxycarbonylpiperazine (3 g) and potassium carbonate (0.82 g) in N,N-dimethylformamide (15 ml) was added octyl bromide (1.87 ml). The mixture was stirred for 10 hours at 70°C. The reaction mixture was added to a mixture of water and ethyl acetate. The organic layer was taken, and dried over magnesium sulfate. The magnesium sulfate was filtered off, and the filtrate was evaporated under reduced pressure. The residue was subjected to column chromatography on silica gel, and eluted with (hexane : ethyl acetate = 9:1). The fractions containing the object compound were combined, and evaporated under reduced pressure to give 1-(4-n-Octyloxyphenyl)-4-tert-butoxycarbonylpiperazine (2.71 g).

IR (KBr) : 1687, 1513, 1241 cm^{-1}

NMR (CDCl_3 , δ) : 0.88 (3H, t, $J=6.2\text{Hz}$), 1.2-1.4 (10H, m), 1.48 (9H, s), 1.65-1.85 (2H, m), 3.00 (4H, t, $J=5.2\text{Hz}$), 3.57 (4H, t, $J=5.2\text{Hz}$), 3.90 (2H, t, $J=6.5\text{Hz}$), 6.83 (2H, dd, $J=6.4$ and 2.1Hz), 6.89 (2H, dd, $J=6.4$ and 2.1Hz)

Preparation 2

A solution of 1-(4-n-Octyloxyphenyl)-4-tert-butoxycarbonylpiperazine (2.61 g) in trifluoroacetic acid (20 ml) was stirred for 4 hours at ambient temperature. The reaction mixture was evaporated under reduced pressure, and to the residue was added a mixture of 1N NaOH aqueous solution and ethyl acetate. The organic layer was taken, and dried over magnesium sulfate. The magnesium sulfate was filtered off, and the filtrate was evaporated under reduced pressure to give 1-(4-n-Octyloxyphenyl)piperazine (0.86 g).

IR (KBr) : 2923, 1513, 1259, 831 cm^{-1}

NMR (CDCl_3 , δ) : 0.88 (3H, t, $J=6.4\text{Hz}$), 1.2-1.53

(10H, m), 1.65-1.85 (2H, m), 3.03 (4H, s), 3.90
(2H, t, J=6.5Hz), 6.83 (2H, dd, J=6.4 and
2.9Hz), 6.90 (2H, dd, J=6.4 and 2.9Hz)

APCI-MASS : m/z = 291 (M⁺+1)

5

Preparation 3

To a suspension of 1-(4-n-Octyloxyphenyl)piperazine
(1 g) and potassium carbonate (0.476 g) in N,N-dimethyl-
formamide (1 ml) was added p-fluorobenzonitrile (0.347 g),
10 and stirred for 5 hours at 160°C. The reaction mixture
was added to a mixture of water and ethyl acetate. The
organic layer was taken, and dried over magnesium sulfate.
The magnesium sulfate was filtered off, and the filtrate
was evaporated under reduced pressure to give 4-[4-(4-n-
15 Octyloxyphenyl)piperazin-1-yl]benzonitrile (0.93 g).

IR (KBr) : 2848, 2217, 1604, 1511, 1241 cm⁻¹

NMR (CDCl₃, δ) : 0.89 (3H, t, J=6.8Hz), 1.2-1.53

(10H, m), 1.65-1.85 (2H, m), 3.20 (4H, t,
J=5.4Hz), 3.48 (4H, t, J=5.4Hz), 3.91 (2H, t,
20 J=6.5Hz), 6.8-7.0 (6H, m), 7.52 (2H, d, J=8.9Hz)

APCI-MASS : m/z = 392 (M⁺+1)

Preparation 4

A mixture of 2,4-Dihydroxybenzaldehyde (5.52 g),
25 potassium carbonate (6.08 g) and octyl bromide (7.73 g) in
acetonitrile (55 ml) was stirred for 16 hours at 60°C.
The solvent of reaction mixture was removed under reduced
pressure, and the residue was dissolved in ethyl acetate,
and washed with water and brine. The separated organic
30 layer was dried over magnesium sulfate. The magnesium
sulfate was filtered off, and the filtrate was evaporated
under reduced pressure. The residue was subjected to
column chromatography on silica gel and eluted with
(hexane : ethyl acetate = 9:1) to give 2-Hydroxy-4-
35 octyloxybenzaldehyde (6.73 g).

NMR (CDCl_3 , δ) : 0.89 (3H, t, $J=8.8\text{Hz}$), 1.2-1.5
(10H, m), 1.8-2.0 (2H, m), 4.0-4.2 (2H, m), 6.42
(1H, s), 6.52 (1H, d, $J=8.7\text{Hz}$), 7.79 (1H, d,
 $J=8.7\text{Hz}$), 10.33 (1H, s)

5 APCI-MASS : $m/z = 257$ (M^++1)

The following compound was obtained according to a
similar manner to that of Preparation 4.

10 Preparation 5

Methyl 3,4-dipentyloxybenzoate

NMR (CDCl_3 , δ) : 0.93 (6H, t, $J=6.0$ and 9.0Hz), 1.3-
2.0 (12H, m), 3.88 (3H, s), 4.04 (4H, m),
6.86 (1H, d, $J=8.4\text{Hz}$), 7.53 (1H, d, $J=2.0\text{Hz}$),
15 7.63 (1H, dd, $J=8.4$ and 2.0Hz)

APCI-MASS : $m/z = 309$ (M^++1)

Preparation 6

A mixture of 4-bromo-4'-pentylbiphenyl (5.04 g),
20 trimethylsilylacetylene (2.4 ml),
tetrakis(triphenylphosphine)palladium (0.96 g),
triphenylphosphine (0.22 g) and cuprous iodide (95 mg) in
piperidine (10 ml) was heated for an hour under
atmospheric pressure of nitrogen at 90°C . The reaction
25 mixture was poured into a mixture of cold water and ethyl
acetate, and adjusted to about pH 1 with 6N hydrochloric
acid. The separated organic layer was washed with water
and brine, and dried over magnesium sulfate. The
magnesium sulfate was filtered off, and the filtrate was
30 evaporated under reduced pressure to give crude 2-[4-(4-
pentylphenyl)phenyl]-1-trimethylsilylacetylene, which was
used for the next reaction without further purification.
Crude mixture was dissolved in a mixture of
dichloromethane (10 ml) and methanol (10 ml), and to the
35 solution was added potassium carbonate (2.75 g) at 0°C .

The mixture was allowed to warm to ambient temperature, and stirred for another 2 hours. The reaction mixture was poured into a mixture of cold water and ethyl acetate, and the resultant precipitate was filtered off. The filtrate was adjusted to about pH 7 with 1N hydrochloric acid, and washed with brine, and dried over magnesium sulfate. The magnesium sulfate was filtered off, and the filtrate was evaporated under reduced pressure to give a crude powder, which was subjected to column chromatography on silica gel (300 ml), and eluted with a mixture of (n-hexane : ethyl acetate = 99:1 - 97:3, V/V) to give 4-(4-Pentylphenyl)phenylacetylene (2.09g).

IR (Nujol) : 3274, 1490 cm^{-1}

NMR (CDCl_3 , δ) : 0.90 (3H, t, $J=6.4\text{Hz}$), 1.30-1.50 (4H, m), 1.50-1.80 (2H, m), 2.64 (2H, t, $J=7.6\text{Hz}$), 7.20-7.30 (2H, m), 7.45-7.60 (6H, m)

APCI-MASS : m/z = 281 (M^++1 + MeOH)

The following compound was obtained according to a similar manner to that of Preparation 6.

Preparation 7

6-Heptyloxynaphthalen-2-yl-acetylene

NMR (CDCl_3 , δ) : 0.90 (3H, t, $J=6.5\text{Hz}$), 1.20-1.60 (8H, m), 1.70-1.90 (2H, m), 3.10 (1H, s), 4.07 (2H, t, $J=6.5\text{Hz}$), 7.08 (1H, d, $J=2.5\text{Hz}$), 7.15 (1H, dd, $J=2.5$ and 8.9Hz), 7.47 (1H, dd, $J=1.6$ and 8.5Hz), 7.64 (1H, d, $J=7.3\text{Hz}$), 7.68 (1H, d, $J=8.5\text{Hz}$), 7.94 (1H, d, $J=1.6\text{Hz}$)

APCI-MASS : m/z = 267 (M^++1)

Preparation 8

To a solution of 4-(4-Pentylphenyl)phenylacetylene (2.09 g) in tetrahydrofuran (30 ml) was added dropwise a solution of lithium diisobutylamide in a mixture of

tetrahydrofuran and n-hexane (1.60 M, 5.6 ml) at -75°C, and the resultant mixture was stirred for an hour at -78°C. To the mixture was added methyl chloroformate (0.72 ml), and the reaction mixture was allowed to warm to ambient temperature. The solution was diluted with ethyl acetate, and washed in turn with water and brine, and dried over magnesium sulfate. The magnesium sulfate was filtered off, and the filtrate was evaporated under reduced pressure to give a crude product, which was subjected to column chromatography on silica gel (150 ml), and eluted with a mixture of (n-hexane : ethyl acetate = 100:0 - 9:1, V/V) to give Methyl 3-[4-(4-pentylphenyl)phenyl]propionate (2.20 g).

IR (Nujol) : 2225, 1712 cm^{-1}

NMR (CDCl_3 , δ) : 0.90 (3H, t, $J=6.5\text{Hz}$), 1.25-1.50 (4H, m), 1.52-1.80 (2H, m), 2.64 (2H, t, $J=7.6\text{Hz}$), 3.85 (3H, s), 7.20-7.35 (2H, m), 7.40-7.70 (6H, m)

APCI-MASS : $m/z = 307$ (M^++1)

The following compound was obtained according to a similar manner to that of Preparation 8.

Preparation 9

Methyl 3-(6-heptyloxynaphthalen-2-yl)propionate

IR (Nujol) : 2219, 1704, 1621 cm^{-1}

NMR (CDCl_3 , δ) : 0.90 (3H, t, $J=6.5\text{Hz}$), 1.20-1.60 (8H, m), 1.70-2.00 (2H, m), 3.86 (3H, s), 4.08 (2H, t, $J=6.5\text{Hz}$), 7.10 (1H, d, $J=2.5\text{Hz}$), 7.17 (1H, dd, $J=2.5$ and 8.9Hz), 7.52 (1H, dd, $J=1.6$ and 8.5Hz), 7.68 (1H, d, $J=7.3\text{Hz}$), 7.72 (1H, d, $J=8.5\text{Hz}$), 8.06 (1H, d, $J=1.6\text{Hz}$)

APCI-MASS : $m/z = 325$ (M^++1)

Preparation 10

A mixture of 4-bromo-4'-pentylbiphenyl (5.0 g), methyl acrylate (2.2 ml), palladium acetate (0.11 g) and tris(o-tolyl)phosphine (0.60 g) in triethylamine (16 ml) was refluxed for 15 hours under nitrogen atmosphere. The reaction mixture was poured into a mixture of cold water and ethyl acetate, and adjusted to about pH 1.5 with 6N hydrochloric acid. The separated organic layer was washed in turn with water and brine, and dried over magnesium sulfate. The magnesium sulfate was filtered off, and the filtrate was evaporated under reduced pressure to give a crude powder, which was subjected to column chromatography on silica gel (200 ml), and eluted with a mixture of (n-hexane : ethyl acetate = 100:0 - 94:6, V/V) to give Methyl 3-[4-(4-pentylphenyl)phenyl]acrylate (4.48 g).

IR (Nujol) : 1718, 1637 cm^{-1}

NMR (CDCl_3 , δ) : 0.91 (3H, t, $J=6.7\text{Hz}$), 1.20-1.50 (4H, m), 1.50-1.80 (2H, m), 2.65 (2H, t, $J=7.4\text{Hz}$), 3.82 (3H, s), 6.47 (1H, d, $J=16.0\text{Hz}$), 7.20-7.35 (2H, m), 7.45-7.68 (6H, m), 7.73 (1H, d, $J=16.0\text{Hz}$)

APCI-MASS : $m/z = 309$ (M^++1)

The following compounds (Preparations 11 to 13) were obtained according to a similar manner to that of Preparation 10.

Preparation 11

Methyl 3-(6-heptyloxynaphthalen-2-yl)acrylate

IR (Nujol) : 1716, 1625, 1459 cm^{-1}

NMR (CDCl_3 , δ) : 0.90 (3H, t, $J=6.5\text{Hz}$), 1.20-1.65 (8H, m), 1.76-1.93 (2H, m), 3.82 (3H, s), 4.07 (2H, t, $J=6.5\text{Hz}$), 6.49 (1H, d, $J=16.0\text{Hz}$), 7.05-7.20 (2H, m), 7.55-7.90 (5H, m)

APCI-MS : $m/z = 327$ (M^++1)

Preparation 12

Methyl 3-[4-(4-heptylphenyl)phenyl]acrylate

NMR (CDCl₃, δ) : 0.88 (3H, t, J=6.5Hz), 1.15-1.50
(8H, m), 1.50-1.75 (2H, m), 2.64 (2H, t,
J=7.6Hz), 3.81 (3H, s), 6.46 (1H, d, J=16.0Hz),
7.26 (2H, d, J=8.2Hz), 7.52 (2H, d, J=8.2Hz),
7.59 (6H, s), 7.73 (1H, d, J=16.0Hz)

APCI-MASS : m/z = 337 (M⁺+1)

10 Preparation 13

Methyl 3-[4-(4-pentyloxyphenyl)phenyl]acrylate

NMR (CDCl₃, δ) : 0.94 (3H, t, J=7.0Hz), 1.30-1.60
(4H, m), 1.70-1.93 (2H, m), 3.82 (3H, s), 4.00
(2H, t, J=6.7Hz), 6.45 (1H, d, J=16.0Hz), 6.90-
7.05 (2H, m), 7.48-8.65 (6H, m), 7.72 (1H, d,
J=16.0Hz)

APCI-MASS : m/z = 325 (M⁺+1)

Preparation 14

20 A mixture of 6-Heptyloxynaphthalen-2-carboxylic acid
(1.00 g) and thionyl chloride (5 ml) was stirredn for 18
hours at ambient temperature, and concentrated under
reduced pressure to give crude 6-heptyloxy-2-naphthoyl
chloride. To a mixture of ethyl isonipecotinate (605 mg),
25 triethylamine (425 mg) and N,N-dimethylaminopyridine (10
mg) in dichloromethane (10 ml) was added crude 6-
heptyloxy-2-naphthoyl chloride, and the mixture was
stirred for 2 hours at ambient temperature, and diluted
with dichloromethane. The mixture was washed with water,
30 1N hydrochloric acid and brine, and dried over magnesium
sulfate. The magnesium sulfate was filtered off, and
filtrate was evaporated under reduced pressure. The
residue was subjected to column chromatography on silica
gel, and eluted with (n-hexane : ethyl acetate = 3:1) to
35 give 4-Ethoxycarbonyl-1-(6-heptyloxy-2-

naphthoyl)piperidine (1.20 g).

NMR (CDCl_3 , δ) : 0.90 (3H, t, $J=6.6\text{Hz}$), 1.2-2.0
(19H, m), 2.5-2.7 (1H, m), 3.0-3.2 (2H, m), 4.1-
4.3 (4H, m), 7.1-7.2 (2H, m), 7.44 (1H, dd,
5 $J=8.4$ and 1.7Hz), 7.72 (1H, d, $J=3.9\text{Hz}$), 7.77
(1H, d, $J=3.9\text{Hz}$), 7.82 (1H, s)

APCI-MASS : $m/z = 426 (M^++1)$

Preparation 15

10 To a mixture of Methyl 3,4-diaminobenzoate (1.91 g)
and triethylamine (0.56 g) in N,N-dimethylformamide (20
ml) was added decanoyl chloride (2.31 g), and the mixture
was stirred for an hour at 0°C . The reaction mixture was
15 diluted with ethyl acetate, and washed with water and
brine. The separated organic layer was dried over
magnesium sulfate. The magnesium sulfate was filtered
off, and filtrate was evaporated under reduced pressure.
The residue was dissolved in methanol (20 ml), and conc.
20 sulfuric acid (0.05 ml) was added, and the mixture was
stirred for 6 hours at 60°C . After cooling, the reaction
mixture was evaporated under reduced pressure. The
residue was diluted with ethyl acetate, and washed with
water and brine. The separated organic layer was dried
25 over magnesium sulfate. The magnesium sulfate was
filtered off, and filtrate was evaporated under reduced
pressure. Purification of the residue by column
chromatography on silica gel eluted with (n-hexane : ethyl
acetate = 3:1) gave

5-Methoxycarbonyl-2-nonylbenzimidazole (1.40 g).

30 IR (KBr pelet) : 2923, 1718, 1623, 1544, 1438, 1413,
1288, 1213, 1085, 750 cm^{-1}

NMR ($\text{DMSO}-d_6$, δ) : 0.84 (3H, t, $J=6.7\text{Hz}$), 1.1-1.4
(12H, m), 1.7-1.9 (2H, m), 2.83 (2H, t,
 $J=7.4\text{Hz}$), 7.56 (1H, d, $J=8.4\text{Hz}$), 7.78 (1H, d,
35 $J=8.4\text{Hz}$), 8.07 (1H, s)

APCI-MASS : $m/z = 303 (M^+ + 1)$

Preparation 16

To a mixture of dimethylmalonate (4 ml), 2-hydroxy-4-octyloxybenzaldehyde (2.50 g) and piperidine (0.1 ml) in methanol (10 ml) was added acetic acid (0.01 ml), and the mixture was stirred for 3 hours at 70°C. The solvents were removed under reduced pressure, and the residue was dissolved in ethyl acetate, and washed with 0.5N hydrochloric acid, water and brine, and dried over magnesium sulfate. The magnesium sulfate was filtered off, and filtrate was evaporated under reduced pressure, and the precipitate was collected by filtration, and washed with n-hexane, and dried to give Methyl 7-octyloxy coumarin-3-carboxylate (0.94 g).

NMR (DMSO- d_6 , δ) : 0.86 (3H, m), 1.2-1.6 (10H, m), 1.7-1.8 (2H, m), 3.81 (3H, s), 4.11 (2H, t, $J=6.4\text{Hz}$), 6.9-7.1 (2H, m), 7.83 (1H, d, $J=9.0\text{Hz}$), 8.75 (1H, s)

APCI-MASS : $m/z = 333 (M^+ + 1)$

Preparation 17

To a mixture of sodium hydride (423 mg) and 4-octylphenol (2.06 g) in tetrahydrofuran (16 ml) was added dropwise ethyl 2-chloroacetoacetate at ambient temperature. The mixture was stirred for 6 hours at 70°C under nitrogen atmosphere, and poured into saturated ammonium chloride aqueous solution. The solution was extracted with ethyl acetate, and the organic layer was washed with water and brine, and dried over magnesium sulfate. The magnesium sulfate was filtered off, and the filtrate was evaporated under reduced pressure. The residue was added to conc. H_2SO_4 (10 ml) at 0°C, and mixture was stirred for 10 minutes. The reaction mixture was poured into ice-water, and adjusted to pH 7.0 with 1N

NaOH aqueous solution, and extracted with ethyl acetate. The organic layer was washed with water and brine, and dried over magnesium sulfate. The magnesium sulfate was filtered off, and the filtrate was evaporated under reduced pressure. The residue was subjected to column-chromatography on silica gel, and eluted with (hexane : ethyl acetate = 95:5). The fractions containing the object compound were combined, and evaporated under reduced pressure to give Ethyl 3-methyl 5-octylbenzo[b]furan-2-carboxylate (1.44 g).

IR (Neat) : 2925, 2854, 1712, 1596, 1463, 1292, 1149, 1089 cm^{-1}

NMR (CDCl_3 , δ) : 0.88 (3H, t, $J=6.7\text{Hz}$), 1.2-1.5 (10H, m), 1.44 (3H, t, $J=7.1\text{Hz}$), 1.6-1.8 (2H, m), 2.58 (3H, s), 2.71 (2H, t, $J=8.0\text{Hz}$), 4.45 (2H, t, $J=7.1\text{Hz}$), 7.2-7.5 (3H, m)

APCI-MASS : $m/z = 317$ (M^++1)

Preparation 18

To a solution of Ethyl 3-amino-4-hydroxybenzoate (1.81 g) and triethylamine (1.53 ml) in dichloromethane (20 ml) was dropwise added decanoyl chloride (2.01 ml) at 0°C . The reaction mixture was stirred for 48 hours at ambient temperature, and washed with water, 0.5N hydrochloric acid, water and brine. The separated organic layer was dried over magnesium sulfate. The magnesium sulfate was filtered off, and the filtrate was evaporated under reduced pressure. To the residue dissolved in xylene (30 ml) was added p-toluene sulfonic acid monohydrate (0.5 g), and the mixture was stirred for 4 hours at 130°C . Ethyl acetate was added to the mixture, and washed with water and brine. The separated organic layer was dried over magnesium sulfate. The magnesium sulfate was filtered off, and the filtrate was evaporated under reduced pressure. Purification of the residue by

column chromatography on silica gel elluted with
(n-hexane : ethyl acetate = 9:1, V/V) gave Ethyl 2-nonyl
benzo[b]oxazole-6-carboxylate (2.36 g).

5 IR (KBr pelet) : 2914, 1722, 1621, 1575, 1470,
1429, 1365, 1290, 1203, 1151, 1115, 1081,
1022 cm^{-1}

NMR (CDCl_3 , δ) : 0.88 (3H, t, $J=6.7\text{Hz}$), 1.2-1.4 (12H,
m), 1.42 (3H, t, $J=7.2\text{Hz}$), 1.90 (2H, m), 2.95
10 (2H, t, $J=7.4\text{Hz}$), 4.40 (2H, q, $J=7.0\text{Hz}$), 7.50
(1H, d, $J=8.5\text{Hz}$), 8.06 (1H, d, $J=8.5\text{Hz}$), 8.37
(1H, s)

APCI-MASS : $m/z = 318$ (M^++1)

Preparation 19

15 A mixture of Methyl 3,4-diaminobenzoate (1.84 g) and
4-hexyloxy benzaldehyde (2.30 g) in nitrobenzene (40 ml)
was stirred for 48 hours at 145°C . After cooling, the
mixture was evaporated under reduced pressure.
Purification of the residue by column chromatography on
20 silica gel eluted with (n-hexane : ethyl acetate = 2:1)
gave 5-Methoxycarbonyl-2-(4-hexyloxyphenyl)benzimidazole
(1.19 g).

NMR (CDCl_3 , δ) : 0.90 (3H, t, $J=6.4\text{Hz}$), 1.2-1.9 (8H,
m), 3.92 (3H, s), 3.90-4.1 (2H, m), 6.93 (2H, d,
25 $J=8.9\text{Hz}$), 7.5-7.8 (1H, br), 7.94 (1H, dd, $J=8.5$
and 1.5Hz), 8.03 (1H, d, $J=8.9\text{Hz}$), 8.2-8.4 (1H,
br)

APCI-MASS : $m/z = 353$ (M^++1)

Preparation 20

30 A mixture of Methyl 3-[4-(4-pentylphenyl)phenyl]-
acrylate (2.0 g) and 10% palladium on carbon (50% wet, 0.2
g) in tetrahydrofuran (20 ml) was stirred for 8 hours
under atmospheric pressure of hydrogen at ambient
35 temparature. The catalyst was filtered off, and the

filtrate was evaporated under reduced pressure to give Methyl 3-[4-(4-pentylphenyl)phenyl]propionate (1.93 g).

NMR (CDCl₃, δ) : 0.90 (3H, t, J=6.8Hz), 1.25-1.50 (4H, m), 1.50-1.75 (2H, m), 2.55-2.75 (4H, m), 2.99 (2H, t, J=8.0Hz), 3.68 (3H, s), 7.10-7.30 (4H, m), 7.40-7.60 (4H, m)

APCI-MASS : m/z = 311 (M⁺+1)

Preparation 21

A mixture of Methyl 3-[4-(4-pentyloxyphenyl)phenyl]-acrylate (2.70 g) and platinum oxide (0.41 g) in tetrahydrofuran (40 ml) was stirred for 8 hours under 3 atom of hydrogen at ambient temperature. The catalyst was filtered off, and the filtrate was evaporated under reduced pressure to give Methyl 3-[4-(4-pentyloxyphenyl)phenyl]propionate (2.70 g).

NMR (CDCl₃, δ) : 0.94 (3H, t, J=7.0Hz), 1.28-1.60 (4H, m), 1.60-1.95 (2H, m), 2.55-2.78 (2H, m), 2.98 (2H, t, J=7.8Hz), 3.98 (2H, t, J=6.5Hz), 6.85-7.05 (2H, m), 7.05-7.30 (2H, m), 7.40-7.55 (4H, m)

APCI-MASS : m/z = 327 (M⁺+1)

The following compound was obtained according to a similar manner to that of Preparation 21.

Preparation 22

Methyl 3-(6-heptyloxynaphthalen-2-yl)propionate

NMR (CDCl₃, δ) : 0.90 (3H, t, J=6.5Hz), 1.20-1.70 (8H, m), 1.70-1.93 (2H, m), 2.70 (2H, t, J=7.7Hz), 3.07 (2H, t, J=7.7Hz), 3.67 (3H, s), 4.05 (2H, t, J=6.5Hz), 7.02-7.20 (2H, m), 7.20-7.38 (2H, m), 7.55 (1H, s), 7.66 (1H, dd, J=3.0 and 8.5Hz)

APCI-MASS : m/z = 329 (M⁺+1)

Preparation 23

To a mixture of Methyl 3-[4-(4-pentylphenyl)phenyl]-acrylate (0.41 g) in tetrahydrofuran (5 ml) was added 3N NaOH aqueous solution (1.3 ml), and the resultant mixture was heated to 85°C for 10 hours. The reaction mixture was poured into a mixture of cold water and ethyl acetate, and adjusted to about pH 2 with 6N hydrochloric acid. The separated organic layer was washed in turn with water and brine, and dried over magnesium sulfate. The magnesium sulfate was filtered off, and the filtrate was evaporated under reduced pressure to give 3-[4-(4-Pentylphenyl)phenyl]acrylic acid (0.41 g).

NMR (DMSO-d₆, δ) : 0.87 (3H, t, J=7.5Hz), 1.15-1.46 (4H, m), 1.48-1.70 (2H, m), 2.61 (2H, t, J=7.4Hz), 6.56 (1H, d, J=16.0Hz), 7.29 (2H, d, J=8.2Hz), 7.60 (2H, d, J=4.0Hz), 7.66 (2H, d, J=4.0Hz), 7.68-7.85 (3H, m)

APCI-MASS : m/z = 295 (M⁺+1)

The following compounds (Preparations 24 to 31) were obtained according to a similar manner to that of Preparation 23.

Preparation 24

3-[4-(4-Pentyloxyphenyl)phenyl]propionic acid

IR (Nujol) : 1697, 1606, 1500 cm⁻¹

NMR (CDCl₃, δ) : 0.94 (3H, t, J=7.1Hz), 1.25-1.60 (4H, m), 1.70-1.95 (2H, m), 2.72 (2H, t, J=7.5Hz), 3.00 (2H, t, J=7.5Hz), 3.99 (2H, t, J=6.5Hz), 6.95 (2H, dd, J=2.1 and 6.7Hz), 7.25 (2H, d, J=8.2Hz), 7.40-7.60 (4H, m)

APCI-MASS : m/z = 313 (M⁺+1)

Preparation 25

3-[4-(4-Heptylphenyl)phenyl]propionic acid

NMR (CDCl₃, δ) : 0.88 (3H, t, J=6.8Hz), 1.15-1.50
(8H, m), 1.50-1.78 (2H, m), 2.65 (2H, t,
J=7.6Hz), 6.48 (1H, d, J=16.0Hz), 7.27 (2H, d,
J=8.2Hz), 7.53 (2H, d, J=8.2Hz), 7.63 (4H, m),
7.83 (1H, d, J=16.0Hz)

APCI-MASS : m/z = 323 (M⁺+1)

Preparation 26

3-[4-(4-Pentylphenyl)phenyl]propionic acid

NMR (CDCl₃, δ) : 0.90 (3H, t, J=6.4Hz), 1.20-1.50
(4H, m), 1.50-1.75 (2H, m), 2.64 (2H, t,
J=8.0Hz), 2.67 (2H, t, J=9.6Hz), 3.00 (2H, t,
J=8.0Hz), 7.15-7.38 (4H, m), 7.38-7.60 (4H, m)

APCI-MASS : m/z = 297 (M⁺+1)

Preparation 27

3-(6-Heptyloxynaphthalen-2-yl)propionic acid

NMR (CDCl₃, δ) : 0.90 (3H, t, J=6.5Hz), 1.20-1.65
(8H, m), 1.75-2.00 (2H, m), 2.75 (2H, t,
J=7.7Hz), 3.09 (2H, t, J=7.7Hz), 4.06 (2H, t,
J=6.5Hz), 7.05-7.15 (2H, m), 7.15-7.35 (2H, m),
7.50-7.73 (2H, m)

APCI-MASS : m/z = 315 (M⁺+1)

Preparation 28

3-(6-Heptyloxynaphthalen-2-yl)acrylic acid

NMR (CDCl₃, δ) : 0.90 (3H, t, J=6.5Hz), 1.15-1.60
(8H, m), 1.75-1.95 (2H, m), 4.09 (2H, t,
J=6.5Hz), 6.51 (1H, d, J=16.0Hz), 7.09-7.30 (2H,
m), 7.65-8.00 (5H, m)

Preparation 29

3-[4-(4-Pentylphenyl)phenyl]propionic acid

NMR (CDCl₃, δ) : 0.91 (3H, t, J=6.5Hz), 1.23-1.50
(4H, m), 1.50-1.80 (2H, m), 2.65 (2H, t,

J=7.6Hz), 7.27 (2H, d, J=8.2Hz), 7.51 (2H, d, J=8.2Hz), 7.58-7.80 (4H, m)

APCI-MASS : m/z = 325 ($M^+ + 1$ + MeOH)

5 Preparation 30

3-(6-Heptyloxynaphthalen-2-yl)propionic acid

IR (Nujol) : 2645, 2198, 1670, 1627 cm^{-1}

NMR (DMSO- d_6 , δ) : 0.85 (3H, t, J=6.5Hz), 1.10-1.60

(8H, m), 1.65-1.90 (2H, m), 4.10 (2H, t,

10 J=6.5Hz), 7.24 (1H, dd, J=2.4 and 8.9Hz), 7.39

(1H, d, J=2.5Hz), 7.55 (1H, dd, J=1.6 and

8.5Hz), 7.8-8.0 (2H, m), 8.22 (1H, d, J=1.6Hz)

APCI-MASS : m/z = 343 ($M^+ + 1$ + MeOH)

15 Preparation 31

4-[5-(4-Pentyloxyphenyl)isoxazolyl-3-yl]benzoic acid

IR (KBr) : 2939, 2867, 1681, 1614, 1429, 1255, 1178,
821 cm^{-1}

NMR (DMSO- d_6 , δ) : 0.91 (3H, t, J=7.1Hz), 1.3-1.5

20 (4H, m), 1.6-1.8 (2H, m), 4.04 (2H, t, J=6.5Hz),

7.11 (2H, d, J=8.9Hz), 7.54 (1H, s), 7.85 (2H,

d, J=8.9Hz), 7.98 (2H, d, J=8.6Hz), 8.11 (2H, d,
J=8.6Hz)

APCI-MASS : m/z = 352 ($M + H$)⁺

25

Preparation 32

To a solution of Ethyl 3-methyl-5-octylbenzo[b]furan-
2-carboxylate (1.44 g) in ethanol (20 ml) was added 10%

NaOH aqueous solution (2.2 ml), and stirred for 2 hours at

30 ambient temperature, and evaporated under reduced

pressure. The residue was adjusted to pH 3.0 with 1N

hydrochloric acid, and extracted with ethyl acetate. The

organic layer was washed with brine, and dried over

magnesium sulfate. The magnesium sulfate was filtered off,

35 and the filtrate was evaporated under reduced pressure to

give 3-Methyl-5-octylbenzo[b]furan-2-carboxylic acid (1.00 g).

IR (KBr pelet) : 2923, 1689, 1664, 1581, 1456, 1319, 1159, 933 cm^{-1}

5 NMR (DMSO-d_6 , δ) : 0.85 (3H, t, $J=6.7\text{Hz}$),
1.2-1.5 (10H, m), 1.5-1.8 (2H, m), 2.49 (3H, s),
2.69 (2H, t, $J=7.9\text{Hz}$), 7.32 (1H, dd, $J=8.5$ and
1.7Hz), 7.52 (1H, d, $J=8.5\text{Hz}$), 7.54 (1H, d,
 $J=1.7\text{Hz}$), 13.2-13.5 (1H, br)

10 APCI-MASS : $m/z = 289$ (M^++1)

The following compounds (Preparations 33 to 39) were obtained according to a similar manner to that of Preparation 32.

15

Preparation 33

3,4-Dipentyloxybenzoic acid

20 NMR (DMSO-d_6 , δ) : 0.89 (6H, t, $J=6.8\text{Hz}$),
1.2-1.5 (8H, m), 1.6-1.8 (4H, m), 3.9-4.1 (4H,
m), 7.02 (1H, d, $J=8.4\text{Hz}$), 7.43 (1H, d,
 $J=1.7\text{Hz}$), 7.53 (1H, dd, $J=8.4$ and 1.7Hz)

APCI-MASS : $m/z = 295$ (M^++1)

Preparation 34

25 1-(6-Heptyloxy-2-naphthoyl)piperidine-4-carboxylic acid

30 NMR (DMSO-d_6 , δ) : 0.88 (3H, t, $J=6.7\text{Hz}$), 1.2-2.0
(14H, m), 2.5-2.6 (1H, m), 2.9-3.2 (2H, br),
3.25 (2H, s), 4.09 (2H, t, $J=6.5\text{Hz}$), 7.20 (1H,
dd, $J=8.9$ and 2.4Hz), 7.36 (1H, d, $J=2.3\text{Hz}$),
7.43 (1H, dd, $J=8.4$ and 1.5Hz), 7.8-8.0 (3H, m),
12.30 (1H, br)

APCI-MASS : $m/z = 398$ (M^++1)

35

Preparation 35

7-Octyloxycoumarin-3-carboxylic acid

IR (KBr) : 1748, 1625, 1558, 1467, 1430, 1386, 1360,
1257, 1217, 1120 cm^{-1}

5 NMR (DMSO- d_6 , δ) : 0.86 (3H, t, $J=6.8\text{Hz}$), 1.2-1.5
(10H, m), 1.6-1.8 (2H, m), 4.11 (2H, t,
 $J=6.4\text{Hz}$), 6.9-7.1 (2H, m), 7.82 (1H, d,
 $J=8.9\text{Hz}$), 8.72 (1H, s), 12.98 (1H, br)
APCI-MASS : $m/z = 319$ (M^++1)

10

Preparation 36

4-(4-Pentyloxyphenyl)cinnamic acid

IR (Nujol) : 2923, 1675, 1500, 1290, 1223, 985,
821 cm^{-1}

15 NMR (DMSO- d_6 , δ) : 0.90 (3H, t, $J=7.0\text{Hz}$), 1.3-1.5
(4H, m), 1.6-1.8 (2H, m), 4.01 (2H, t, $J=6.5\text{Hz}$),
6.54 (1H, d, $J=16.0\text{Hz}$), 7.02 (2H, d, $J=8.8\text{Hz}$),
7.5-7.8 (7H, m)
APCI-MASS : $m/z = 311$ (M^++1)

20

Preparation 37

2-Nonylbenzoxazole-6-carboxylic acid

25 NMR (DMSO- d_6 , δ) : 0.84 (3H, t, $J=6.7\text{Hz}$), 1.2-1.5
(12H, m), 1.7-1.9 (2H, m), 2.96 (2H, t,
 $J=7.4\text{Hz}$), 7.76 (1H, d, $J=8.4\text{Hz}$), 7.98 (1H, d,
 $J=8.4\text{Hz}$), 8.19 (1H, s)
APCI-MASS : $m/z = 290$ (M^++1)

Preparation 38

30 2-(4-Hexyloxyphenyl)benzimidazole-5-carboxylic acid
NMR (DMSO- d_6 , δ) : 0.8-1.0 (3H, m), 1.3-1.6 (6H, m),
1.7-1.8 (2H, m), 4.06 (2H, t, $J=6.4\text{Hz}$), 7.12
(2H, d, $J=8.8\text{Hz}$), 7.6-7.9 (2H, m), 8.1-8.2 (3H,
m), 13.00 (1H, br)
35 APCI-MASS : $m/z = 339$ (M^++1)

Preparation 39

2-Nonylbenzimidazole-5-carboxylic acid

NMR (DMSO-d₆, δ) : 0.85 (3H, t, J=6.7Hz), 1.1-1.4
(12H, m), 2.7-2.9 (2H, m), 2.96 (2H, t,
J=7.6Hz), 3.6-5.2 (1H, br), 7.66 (1H, d,
J=8.4Hz), 7.90 (1H, d, J=8.4Hz), 8.15 (1H, s)

APCI-MASS : m/z = 289 (M⁺+1)

Preparation 40

A solution of 4-[4-(4-Octyloxyphenyl)piperazin-1-yl]benzonitrile (0.5 g) in 20% H₂SO₄ aqueous solution (30 ml) and acetic acid (20 ml) was refluxed for 9 hours. The reaction mixture was pulverized with water. The precipitate was collected by filtration, and added to a mixture of water, tetrahydrofuran and ethyl acetate, and adjusted to pH 2.5 with 1N NaOH aqueous solution. The organic layer was taken, and dried over magnesium sulfate. The magnesium sulfate was filtered off, and the filtrate was evaporated under reduced pressure to give 4-[4-(4-Octyloxyphenyl)piperazin-1-yl]benzoic acid (388 mg).

IR (KBr) : 2929, 1664, 1600, 1510, 1240 cm⁻¹

NMR (DMSO-d₆, δ) : 0.86 (3H, t, J=6.6Hz), 1.2-1.5
(10H, m), 1.5-1.8 (2H, m), 3.13 (4H, t,
J=5.3Hz), 3.44 (4H, t, J=5.3Hz), 3.88 (2H, t,
J=6.5Hz), 6.83 (2H, d, J=9.2Hz), 6.94 (2H, d,
J=9.2Hz), 7.02 (2H, d, J=9.0Hz), 7.79 (2H, d,
J=9.0Hz)

APCI-MASS : m/z = 411 (M⁺+1)

Preparation 41

To a suspension of sodium hydride (60% suspension in mineral oil) (0.296 g) in N,N-dimethylformamide (14 ml) was added 1,2,4-triazole (0.511 g) and 4-[4-(8-bromooctyloxy)phenyl]benzoic acid (1 g), and was stirred for 5 hours at 120°C. The reaction mixture was added to a

mixture of water and ethyl acetate, and adjusted to pH 2.5 with conc. hydrochloric acid. The organic layer was taken and dried over magnesium sulfate. The magnesium sulfate was filtered off, and the filtrate was evaporated under reduced pressure to give 4-[4-[8-(1,2,4-Triazol-1-yl)octyloxy]phenyl]benzoic acid (0.81 g).

IR (KBr) : 2940, 1689, 1604, 1297, 1189 cm^{-1}

NMR (DMSO- d_6 , δ) : 1.1-1.53 (8H, m), 1.6-1.9 (4H, m), 4.00 (2H, t, $J=6.3\text{Hz}$), 4.16 (2H, t, $J=7.0\text{Hz}$), 7.03 (2H, d, $J=8.7\text{Hz}$), 7.67 (2H, d, $J=8.7\text{Hz}$), 7.75 (2H, d, $J=8.4\text{Hz}$), 7.95 (1H, s), 7.99 (2H, d, $J=8.4\text{Hz}$), 8.51 (1H, s), 12.9 (1H, s)

APCI-MASS : $m/z = 394$ ($M^+ + 1$)

Preparation 42

A mixture of 2-Carbamoyl-5-methoxybenzo[b]thiophene (2.0 g), acetic acid (5 ml) and 48% hydrobromic acid (20 ml) was stirred for 16 hours at 110°C , and the mixture was poured into the ice-water. The resulting precipitate was collected by filtration, and dried to give 5-Hydroxybenzo[b]thiophene-2-carboxylic acid (1.66 g).

NMR (DMSO- d_6 , δ) : 7.03 (1H, dd, $J=8.8$ and 0.6Hz), 7.31 (1H, d, $J=0.6\text{Hz}$), 7.81 (1H, d, $J=8.8\text{Hz}$), 7.96 (1H, s), 9.64 (1H, s), 13.32 (1H, s)

APCI-MASS : $m/z = 195$ ($M^+ + 1$)

Preparation 43

A solution of (S)-2-Tert-butoxycarbonyl-1,2,3,4-tetrahydro-7-hydroxyisoquinoline-3-carboxylic acid (1 g) in a mixture of 10% NaOH aqueous solution (2.73 ml) and dimethylsulfoxide (11 ml) was stirred for half an hour at 80°C . Then, octyl bromide (0.589 ml) was added thereto, and stirred for 4 hours at 60°C . The reaction mixture was added to a mixture of water and ethyl acetate, and

adjusted to pH 2.5 with conc. hydrochloric acid. The organic layer was taken, and dried over magnesium sulfate. The magnesium sulfate was filtered off, and the filtrate was evaporated under reduced pressure to give (S)-2-Tert-butoxycarbonyl-1,2,3,4-tetrahydro-7-octyloxyisoquinoline-3-carboxylic acid (1.30 g).

IR (Neat) : 2929, 1743, 1704, 1164 cm^{-1}

NMR (CDCl_3 , δ) : 0.89 (3H, t, $J=6.1\text{Hz}$), 1.1-1.6 (10H, m), 1.41 + 1.51 (9H, s, cis + trans), 1.75 (2H, quint, $J=6.5\text{Hz}$), 3.10 (2H, m), 3.90 (2H, t, $J=3.9\text{Hz}$), 4.42 (1H, d, $J=16.8\text{Hz}$), 4.65 (1H, d, $J=16.8\text{Hz}$), 4.74 + 5.09 (1H, m, cis + trans), 6.5-6.8 (2H, m), 7.03 (1H, d, $J=8.3\text{Hz}$)

APCI-MASS : m/z = 306 ($M^+ + 1$ -Boc)

The following compounds (Preparations 44 to 45) were obtained according to a similar manner to that of Preparation 43.

Preparation 44

5-Octyloxybenzo[b]thiophene-2-carboxylic acid

IR (KBr) : 1673, 1666, 1600, 1517, 1409, 1267, 1214, 1153, 865 cm^{-1}

NMR ($\text{DMSO}-d_6$, δ) : 0.86 (3H, t, $J=6.7\text{Hz}$), 1.2-1.5 (10H, m), 1.7-1.9 (2H, m), 4.02 (2H, t, $J=6.4\text{Hz}$), 7.13 (1H, dd, $J=8.9$ and 0.6Hz), 7.51 (1H, d, $J=0.6\text{Hz}$), 7.90 (1H, d, $J=9.0\text{Hz}$), 7.99 (1H, s)

APCI-MASS : m/z = 307 ($M^+ + 1$)

Preparation 45

4-[4-(4-Hexyloxyphenyl)piperazin-1-yl]benzoic acid dihydrochloride

IR (KBr) : 1668, 1600, 1510, 1228 cm^{-1}

NMR ($\text{DMSO}-d_6$, δ) : 0.88 (3H, t, $J=6.9\text{Hz}$), 1.2-1.5

(6H, m), 1.6-1.9 (2H, m), 3.0-3.2 (4H, m), 3.3-3.5 (4H, m), 3.88 (2H, t, J=6.3Hz), 6.83 (2H, d, J=9Hz), 6.9-7.1 (4H, m), 7.79 (2H, d, J=8.8Hz), 12.32 (1H, s)

5 APCI-MASS : m/z = 383 (M+H⁺)

Preparation 46

To a suspension of dimethyl terephthalate (1.94 g) and potassium t-butoxide (2.24 g) in tetrahydrofuran (30 ml) was added 4-pentyloxyacetophenone (1.59 g) in tetrahydrofuran (10 ml) at 70°C dropwise. The mixture was refluxed for 30 minutes and poured into 1N HCl (50 ml). The mixture was extracted with ethyl acetate (100 ml) and the organic layer was washed with H₂O (100 ml), brine (100 ml) and evaporated under reduced pressure. The residue was triturated with acetonitrile (20 ml), collected by filtration and dried under reduced pressure to give 1-(4-Methoxycarbonylphenyl)-3-(4-pentyloxyphenyl)propane-1,3-dione (2.41 g) as yellow solid.

20 IR (KBr) : 3475, 2956, 2923, 1720, 1606, 1508, 1284, 1176, 1108, 769 cm⁻¹

NMR (CDCl₃, δ) : 0.95 (3H, t, J=7.0Hz), 1.3-1.5 (4H, m), 1.7-2.0 (2H, m), 3.96 (3H, s), 4.04 (2H, t, J=6.5Hz), 6.82 (1H, s), 6.96 (2H, d, J=8.9Hz), 8.0-8.1 (4H, m), 8.14 (2H, m, J=8.7Hz), 12-13 (1H, br)

25 APCI-MASS : m/z = 369 (M+H⁺)

Preparation 47

30 The solution of 1-(4-Methoxycarbonylphenyl)-3-(4-pentyloxyphenyl)propane-1,3-dione (1.00 g) and hydroxylamine hydrochloride (567 mg) in methanol (10 ml) was refluxed for 10 hours. The reaction mixture was diluted with ethyl acetate (50 ml) and washed with water (50 ml x 2), brine (50 ml). The organic layer was dried

over magnesium sulfate and the solvents were removed under reduced pressure. The residue was triturated with acetonitrile (10 ml), collected by filtration, and dried under reduced pressure to give Methyl 4-[5-(4-pentyloxyphenyl)isoxazol-3-yl]benzoate (0.74 g).

IR (KBr) : 2942, 2873, 1716, 1616, 1508, 1280, 1108 cm^{-1}

NMR (CDCl_3 , δ) : 0.95 (3H, t, $J=6.9\text{Hz}$), 1.3-1.6 (4H, m), 1.8-2.0 (2H, m), 3.95 (3H, s), 4.02 (2H, t, $J=6.5\text{Hz}$), 6.74 (1H, s), 6.99 (2H, d, $J=8.8\text{Hz}$), 7.76 (2H, d, $J=8.8\text{Hz}$), 7.93 (2H, d, $J=8.5\text{Hz}$), 8.14 (2H, d, $J=8.5\text{Hz}$)

APCI-MASS : $m/z = 366$ ($M+H$)⁺

15 Preparation 48

A solution of 4-[4-(8-Bromooctyloxy)phenyl]benzoic acid (1 g) in a mixture of sodium methylate (28% solution in methanol) (10 ml) and N,N-dimethylformamide (5 ml) was refluxed for 5 hours. The reaction mixture was added to a mixture of water and ethyl acetate and adjusted to pH 2.0 with conc. HCl. The organic layer was taken and dried over magnesium sulfate. The magnesium sulfate was filtered off, and the filtrate was evaporated under reduced pressure to give 4-[4-(8-Methoxyoctyloxy)phenyl]benzoic acid (0.77 g).

IR (KBr) : 2935, 1685, 835, 773 cm^{-1}

NMR (CDCl_3 , δ) : 1.27-1.7 (10H, m), 1.7-1.95 (2H, m), 3.34 (3H, s), 3.38 (2H, t, $J=6.4\text{Hz}$), 4.01 (2H, t, $J=6.5\text{Hz}$), 6.99 (2H, d, $J=8.7\text{Hz}$), 7.58 (2H, d, $J=8.7\text{Hz}$), 7.66 (2H, d, $J=8.4\text{Hz}$), 8.15 (2H, d, $J=8.4\text{Hz}$)

APCI-MASS : $m/z = 339$ ($M^+ + H - H_2O$)

Preparation 49

35 To a suspension of 1-Hydroxybenzotriazole (0.283 g)

and 6-octyloxymethylpicolinic acid (0.505 g) in dichloromethane (15 ml) was added 1-ethyl-3-(3'-dimethylaminopropyl)carbodiimide hydrochloride (WSCD·HCl) (0.473 g), and stirred for 3 hours at ambient temperature.

5 The reaction mixture was poured into water. The organic layer was taken, and dried over magnesium sulfate. The magnesium sulfate was filtered off, and the filtrate was evaporated under reduced pressure to give 1-(6-Octyloxymethylpicolinoyl)benzotriazole 3-oxide (737 mg).

10 IR (Neat) : 1793, 1654, 1591, 1039 cm^{-1}

The following compounds [Preparations 50 to 66] were obtained according to a similar manner to that of Preparation 49.

15

Preparation 50

1-[4-(4-Octyloxyphenyl)piperazin-1-yl]benzoyl]-benzotriazole 3-oxide

IR (KBr) : 1783, 1600, 1511, 1232, 1184 cm^{-1}

20

NMR (CDCl_3 , δ) : 0.89 (3H, t, $J=6.6\text{Hz}$), 1.2-1.65 (10H, m), 1.65-1.9 (2H, m), 3.24 (4H, t, $J=5.3\text{Hz}$), 3.62 (4H, t, $J=5.3\text{Hz}$), 3.93 (2H, t, $J=6.5\text{Hz}$), 6.8-7.1 (6H, m), 7.35-7.63 (3H, m), 8.0-8.25 (3H, m)

25

Preparation 51

1-[4-[4-[8-(1,2,4-Triazol-1-yl)octyloxy]phenyl]-benzoyl]benzotriazole 3-oxide

IR (KBr) : 1776, 1600, 1193, 983 cm^{-1}

30

NMR (CDCl_3 , δ) : 1.2-2.0 (12H, m), 4.03 (2H, t, $J=6.4\text{Hz}$), 4.18 (2H, t, $J=7.1\text{Hz}$), 7.02 (2H, d, $J=8.7\text{Hz}$), 7.4-7.63 (3H, m), 7.63 (2H, d, $J=8.7\text{Hz}$), 7.79 (2H, d, $J=8.3\text{Hz}$), 7.95 (1H, s), 8.06 (1H, s), 8.12 (1H, d, $J=7.7\text{Hz}$), 8.32 (2H, d, $J=8.3\text{Hz}$)

35

APCI-MASS : $m/z = 511$ ($M^+ + 1$)

Preparation 52

1-[2-Methyl-2-(4-octyloxyphenoxy)propionyl]-
5 benzotriazole 3-oxide

IR (Neat) : 2927, 1810, 1504, 1047 cm^{-1}

Preparation 53

1-[2-(4-Octyloxyphenoxy)propionyl]benzotriazole
10 3-oxide

IR (KBr) : 2954, 1812, 1513, 1232 cm^{-1}

Preparation 54

1-[(S)-2-tert-Butoxycarbonyl-1,2,3,4-tetrahydro-7-
15 octyloxyisoquinolin-3-yl-carbonyl]benzotriazole 3-oxide

IR (Neat) : 2929, 1816, 1739, 1704, 1392 cm^{-1}

Preparation 55

Succinimido 4-(4-n-octyloxyphenyl)piperazine-1-
20 carboxylate

IR (KBr) : 2925, 1758, 1743, 1513, 1241 cm^{-1}

NMR (CDCl_3 , δ) : 0.89 (3H, t, $J=6.8\text{Hz}$), 1.2-1.5

(10H, m), 1.65-1.85 (2H, m), 2.83 (4H, s),

3.0-3.2 (2H, m), 3.6-3.85 (2H, m), 3.91 (2H, t,

25 $J=6.5\text{Hz}$), 6.84 (2H, dd, $J=8.5$ and 2.7Hz), 6.90

(2H, dd, $J=8.5$ and 2.7Hz)

APCI-MASS : $m/z = 432$ ($M^+ + 1$)

Preparation 56

(6-Heptyloxy-2-naphthyl)methylsuccinimido carbonate

IR (KBr) : 1878, 1832, 1787, 1735, 1209 cm^{-1}

NMR (CDCl_3 , δ) : 0.90 (3H, t, $J=6.2\text{Hz}$), 1.2-1.6 (8H,

m), 1.73-2.0 (2H, m), 2.83 (4H, s), 4.07 (2H, t,

$J=6.5\text{Hz}$), 5.44 (2H, s), 7.13 (1H, d, $J=2.4\text{Hz}$),

35 7.17 (1H, dd, $J=8.8$ and 2.4Hz), 7.44 (1H, dd,

J=8.4 and 1.6Hz), 7.67-7.85 (3H, m)

Preparation 57

1-(3,4-Dipentyloxybenzoyl)benzotriazole 3-oxide

5 IR (KBr) : 2952, 1774, 1594, 1515, 1430, 1272, 1147,
1089 cm^{-1}

NMR (CDCl_3 , δ) : 0.9-1.1 (6H, m), 1.3-1.6 (8H, m),
1.8-2.1 (4H, m), 4.0-4.2 (4H, m), 6.99 (1H, d,
J=8.5Hz), 7.4-7.6 (3H, m), 7.68 (1H, d,
10 J=2.0Hz), 7.92 (1H, dd, J=8.5 and 2.0Hz), 8.10
(1H, d, J=8.5Hz)

APCI-MASS : m/z = 412 (M^+ +1)

Preparation 58

15 1-(7-Octyloxycoumarin-3-yl-carbonyl)benzotriazole
3-oxide

IR (KBr) : 2925, 1754, 1716, 1610, 1548, 1282, 1199,
1172, 1139, 1064, 781, 750 cm^{-1}

20 NMR ($\text{DMSO}-d_6$, δ) : 0.86 (3H, t, J=7.8Hz), 1.2-1.5
(10H, m), 1.6-1.8 (2H, m), 4.11 (2H, t,
J=6.5Hz), 6.9-7.1 (2H, m), 7.41 (1H, t,
J=7.2Hz), 7.54 (1H, t, J=7.2Hz), 7.72 (1H, d,
J=8.3Hz), 7.82 (1H, d, J=8.3Hz), 7.99 (1H, d,
J=8.3Hz), 8.72 (1H, s)

25 APCI-MASS : m/z = 436 (M^+ +1)

Preparation 59

1-[4-(4-Pentyloxyphenyl)cinnamoyl]benzotriazole 3-
oxide

30 IR (Nujol) : 2854, 1778, 1708, 1620, 1597, 1494,
1459, 1434, 1377, 1350, 1250, 1188, 1138, 1086,
978 cm^{-1}

Preparation 60

35 1-(5-Octyloxybenzo[b]thiophen-2-yl-carbonyl)-

benzotriazole 3-oxide

IR (KBr) : 2950, 1776, 1517, 1342, 1211, 1151 cm^{-1}

NMR (DMSO-d_6 , δ) : 0.86 (3H, t, $J=6.7\text{Hz}$), 1.2-1.5
(10H, m), 1.7-1.9 (2H, m), 4.01 (2H, t,
5 $J=6.4\text{Hz}$), 7.13 (1H, dd, $J=8.8$ and 2.4Hz), 7.42
(1H, d, $J=7.1\text{Hz}$), 7.5-7.6 (3H, m), 7.72 (1H, d,
 $J=8.4\text{Hz}$), 7.89 (1H, d, $J=8.8\text{Hz}$), 7.9-8.1 (2H, m)

APCI-MASS : $m/z = 424$ (M^++1)

10 Preparation 61

1-(3-Methyl-5-octylbenzo[b]furan-2-yl-carbonyl)-
benzotriazole 3-oxide

IR (KBr) : 1776, 1575, 1469, 1363, 1324, 1276, 1114,
1027 cm^{-1}

15 NMR (CDCl_3 , δ) : 0.89 (3H, t, $J=6.7\text{Hz}$), 1.2-1.5
(10H, m), 2.6-2.8 (2H, m), 2.71 (3H, s), 2.76
(2H, t, $J=7.4\text{Hz}$), 7.4-7.6 (6H, m), 8.12 (1H, s)

APCI-MASS : $m/z = 406$ (M^++1)

20 Preparation 62

1-(2-Nonylbenzoxazol-5-yl-carbonyl)benzotriazole
3-oxide

IR (KBr) : 2980, 1783, 1623, 1573, 1276, 1151, 1091,
989 cm^{-1}

25 NMR (DMSO-d_6 , δ) : 0.84 (3H, t, $J=6.8\text{Hz}$), 1.1-1.4
(12H, m), 1.81 (2H, t, $J=7.2\text{Hz}$), 2.96 (3H, t,
 $J=7.4\text{Hz}$), 7.41 (1H, t, $J=7.0\text{Hz}$), 7.54 (1H, t,
 $J=7.0\text{Hz}$), 7.74 (2H, t, $J=7.0\text{Hz}$), 7.98 (2H, d,
 $J=7.0\text{Hz}$), 8.19 (1H, s)

30 APCI-MASS : $m/z = 407$ (M^++1)

Preparation 63

1-[2-(4-Hexyloxyphenyl)benzimidazol-5-yl-carbonyl]-
benzotriazole 3-oxide

35 IR (KBr) : 3160, 2931, 2863, 1778, 1612, 1502, 1448,

000072.0519
20250622

1388, 1294, 1247, 1174, 1097, 1010, 732 cm^{-1}
NMR (DMSO-d_6 , δ) : 0.89 (3H, t, $J=6.7\text{Hz}$), 1.2-1.5
(6H, m), 1.7-1.8 (2H, m), 4.08 (2H, t, $J=6.4\text{Hz}$),
7.16 (2H, d, $J=8.7\text{Hz}$), 7.6-8.4 (9H, m), 8.3-8.6
(1H, br)
APCI-MASS : $m/z = 456$ (M^++1)

Preparation 64

1-[4-[4-(8-Methoxyoctyloxy)phenyl]benzoyl]-
benzotriazole-3-oxide
IR (KBr) : 2931, 1793, 1770, 1600 cm^{-1}
NMR (CDCl_3 , δ) : 1.2-1.7 (10H, m), 1.7-1.93 (2H, m),
3.34 (3H, s), 3.38 (2H, t, $J=6.4\text{Hz}$), 4.03 (2H,
t, $J=6.5\text{Hz}$), 7.03 (2H, d, $J=8.8\text{Hz}$), 7.4-7.7 (3H,
m), 7.63 (2H, d, $J=8.8\text{Hz}$), 7.79 (2H, d,
 $J=8.6\text{Hz}$), 8.12 (1H, d, $J=8.2\text{Hz}$), 8.32 (2H, d,
 $J=8.6\text{Hz}$)

Preparation 65

1-[4-[4-(4-Hexyloxyphenyl)piperazin-1-
yl]benzoyl]benzotriazole 3-oxide
IR (KBr) : 1770, 1604, 1510, 1232, 1186 cm^{-1}
NMR (CDCl_3 , δ) : 0.91 (3H, t, $J=6.6\text{Hz}$), 1.2-1.6 (6H,
m), 1.6-1.9 (2H, m), 3.1-3.3 (4H, m), 3.5-3.7
(4H, m), 3.93 (2H, t, $J=6.5\text{Hz}$), 6.87 (2H, d,
 $J=9.2\text{Hz}$), 6.96 (2H, d, $J=9.2\text{Hz}$), 7.00 (2H, d,
 $J=9.0\text{Hz}$), 7.3-7.7 (3H, m), 8.10 (1H, d,
 $J=8.2\text{Hz}$), 8.15 (2H, d, $J=9.0\text{Hz}$)
APCI-MASS : $m/z = 500$ ($M+H^+$)

Preparation 66

1-[4-[5-(4-Pentyloxyphenyl)isoxazol-3-yl]benzoyl]-
benzotriazole 3-oxide
IR (KBr) : 2950, 2837, 1774, 1616, 1508, 1452, 1251,
1006 cm^{-1}

NMR (CDCl₃, δ) : 0.95 (3H, t, J=7.1Hz), 1.3-1.5 (4H, m), 1.8-2.0 (2H, m), 4.04 (2H, t, J=6.5Hz), 6.81 (1H, s), 7.0-7.1 (3H, m), 7.4-7.6 (3H, m), 7.80 (2H, d, J=8.8Hz), 8.0-8.2 (3H, m), 8.40 (2H, d, J=8.4Hz)

APCI-MASS : m/z = 469 (M+H)⁺

Preparation 67

To a suspension of 1-hydroxybenzotriazole (0.20 g) and 4-(4-pentylphenyl)cinnamic acid (0.40 g) in dichloromethane (12.0 ml) was added 1-ethyl-3-(3'-dimethylaminopropyl)carbodiimide hydrochloride (0.33 g) (WSCD·HCl), and the mixture was stirred for 12 hours at ambient temperature. The reaction mixture was diluted with dichloromethane, and washed with brine, and dried over magnesium sulfate. After magnesium sulfate was filtered off, evaporation of the filtrate and trituration with acetonitrile gave 1-[4-(4-Pentylphenyl)cinnamoyl]benzotriazole 3-oxide (0.24 g).

NMR (CDCl₃, δ) : 0.91 (3H, t, J=6.6Hz), 1.20-1.50 (4H, m), 1.50-1.75 (2H, m), 2.66 (2H, t, J=8.0Hz), 7.20-8.25 (11H, m), 8.55 (1H, d, J=8.4Hz)

APCI-MASS : m/z = 412 (M⁺+1)

The following compounds (Preparations 68 to 73) were obtained according to a similar manner to that of Preparation 67.

Preparation 68

1-[3-[4-(4-Pentyloxyphenyl)phenyl]-2-propanoyl]-benzotriazole 3-oxide

NMR (CDCl₃, δ) : 0.90-1.05 (3H, m), 1.30-1.65 (4H, m), 1.70-1.95 (2H, m), 3.10-3.60 (4H, m), 3.90-4.10 (2H, m), 6.88-7.08 (2H, m),

7.20-8.50 (10H, m)

APCI-MASS : $m/z = 430 (M^+ + 1)$

Preparation 69

5 1-[4-(4-Heptylphenyl)cinnamoyl]benzotriazole 3-oxide

NMR ($CDCl_3$, δ) : 0.89 (3H, t, $J=6.7\text{Hz}$), 1.20-1.50

(8H, m), 1.50-1.80 (2H, m), 2.66 (2H, t,

$J=7.6\text{Hz}$), 6.70-8.60 (12H, m)

APCI-MASS : $m/z = 440 (M^+ + 1)$

10

Preparation 70

1-[3-[4-(4-Pentylphenyl)phenyl]-2-propanoyl]-
benzotriazole 3-oxide

NMR ($CDCl_3$, δ) : 0.90 (3H, t, $J=6.8\text{Hz}$), 1.20-1.50

15 (4H, m), 1.50-1.76 (2H, m), 2.63 (2H, t,

$J=7.4\text{Hz}$), 3.21 (2H, t, $J=7.3\text{Hz}$), 3.51 (2H, t,

$J=7.3\text{Hz}$), 7.20-7.45 (4H, m), 7.45-7.70 (5H, m),

7.78 (1H, dt, $J=1.0$ and 7.2Hz), 8.00 (1H, d,

$J=8.2\text{Hz}$), 8.42 (1H, d, $J=8.4\text{Hz}$)

20 APCI-MASS : $m/z = 414 (M^+ + 1)$

Preparation 71

1-[3-(6-Heptyloxynaphthalen-2-yl)propanoyl]-
benzotriazole 3-oxide

25 NMR ($CDCl_3$, δ) : 0.80-1.10 (3H, m), 1.20-1.70 (8H,
m), 1.70-2.00 (2H, m), 3.10-3.70 (4H, m), 4.00-
4.18 (2H, m), 6.80-8.50 (10H, m)

APCI-MASS : $m/z = 432 (M^+ + 1)$

30 Preparation 72

1-[3-(6-Heptyloxynaphthalen-2-yl)propenoyl]-
benzotriazole 3-oxide

NMR ($CDCl_3$, δ) : 0.90 (3H, t, $J=6.5\text{Hz}$), 1.20-1.65

(8H, m), 1.75-1.95 (2H, m), 4.10 (2H, d,

35 $J=6.5\text{Hz}$), 6.75-8.62 (8H, m)

APCI-MASS : $m/z = 430 (M^+ + 1)$

Preparation 73

1-(4-Hexylphenylbenzoyl)benzotriazole 3-oxide

5 NMR ($CDCl_3$, δ) : 0.90 (3H, t, $J=4.4\text{Hz}$), 1.2-1.5 (6H, m), 1.6-1.8 (2H, m), 2.68 (2H, t, $J=8.0\text{Hz}$), 7.32 (2H, d, $J=8.2\text{Hz}$), 7.4-7.7 (5H, m), 7.81 (2H, d, $J=6.6\text{Hz}$), 8.10 (2H, d, $J=8.1\text{Hz}$), 8.32 (2H, d, $J=7.6\text{Hz}$)

10 APCI-MASS : $m/z = 400 (M^+ + 1)$

Preparation 74

To a solution of 4-octyloxyphenol (1 g) in dimethylformamide (10 ml) and pyridine (0.364 ml) was added
15 N,N'-disuccinimidylcarbonate (1.16 g). The mixture was stirred for 12 hours at ambient temperature. The reaction mixture was added to a mixture of water and ethyl acetate. The organic layer was taken, and dried over magnesium sulfate. The magnesium sulfate was filtered off, and the
20 filtrate was evaporated under reduced pressure to give 4-Octyloxyphenylsuccinimidyl carbonate (0.59 g).

IR (KBr) : 2927, 1876, 1832, 1735 cm^{-1}

25 NMR ($CDCl_3$, δ) : 0.89 (3H, t, $J=6.3\text{Hz}$), 1.2-1.55 (10H, m), 1.67-1.87 (2H, m), 2.87 (4H, s), 3.94 (2H, t, $J=6.5\text{Hz}$), 6.89 (2H, d, $J=9.2\text{Hz}$), 7.17 (2H, d, $J=9.2\text{Hz}$)

APCI-MASS : $m/z = 364 (M^+ + 1)$

30 The following compounds (Preparations 75 to 88) were obtained according to a similar manner to that of Preparation 1.

Preparation 75

Methyl 4-[4-(6-phenylpyridazin-3-yl-oxy)phenyl]benzoate

35 IR (KBr) : 1708, 1427, 1280, 1187, 1112 cm^{-1}

NMR (CDCl₃, δ) : 3.95 (3H, s), 7.2-7.7 (10H, m),
7.92 (1H, d, J=9.2Hz), 8.0-8.2 (4H, m)
APCI-MASS : m/z = 383 (M+H)⁺

5 Preparation 76

Methyl 4-[4-(5-bromopentyloxy)phenyl]benzoate

IR (KBr) : 2946, 2871, 1716, 1602, 1294, 1199, 1112,
837 cm⁻¹

10 NMR (CDCl₃, δ) : 1.7-2.0 (6H, m), 3.45 (2H, t,
J=6.7Hz), 3.93 (3H, s), 4.02 (2H, t, J=6.1Hz), 6.97
(2H, d, J=8.7Hz), 7.56 (2H, d, J=8.7Hz), 7.61 (2H,
d, J=8.3Hz), 8.07 (2H, d, J=8.3Hz)
APCI-MASS : m/z = 378 (M+H)⁺

15 Preparation 77

Methyl 4-[4-(5-phenoxy-pentyloxy)phenyl]benzoate

IR (KBr) : 2944, 2931, 1720, 1600, 1492, 1197,
1110 cm⁻¹

20 NMR (CDCl₃, δ) : 1.6-1.8 (2H, m), 1.8-2.0 (4H, m),
3.93 (3H, s), 4.00 (2H, t, J=6.3Hz), 4.04 (2H, t,
J=6.3Hz), 6.9-7.1 (5H, m), 7.3-7.4 (2H, m), 7.56
(2H, d, J=8.7Hz), 7.62 (2H, d, J=8.3Hz), 8.07 (2H,
d, J=8.3Hz)
APCI-MASS : m/z = 391 (M+H)⁺

25

Preparation 78

1-[2-(4-Cyclohexylphenylamino)ethyl]-2-oxazolidone
hydrochloride

IR (KBr) : 2923.6, 2852.2, 1747.2, 1683.6 cm⁻¹

30 NMR (DMSO-d₆, δ) : 1.1-1.5 (6H, m), 1.6-1.9 (4H, m),
2.3-2.6 (1H, m), 3.3-3.5 (4H, m), 3.58 (2H, dd,
J=9.4 and 7.4Hz), 4.22 (2H, dd, J=9.4 and 7.4Hz),
7.1-7.4 (4H, m)

35 Preparation 79

Methyl 4-[4-(8-hydroxyoctyloxy)phenyl]benzoate

IR (KBr) : 3250, 2933, 2856, 1724, 1602, 1436, 1292,
1199 cm^{-1}

5 NMR (CDCl_3 , δ) : 1.3-1.9 (12H, m), 3.6-3.8 (2H, br),
3.93 (3H, s), 4.00 (2H, t, $J=6.7\text{Hz}$), 4.82 (1H, s),
7.68 (2H, d, $J=8.7\text{Hz}$), 7.56 (2H, d, $J=8.7\text{Hz}$), 7.62
(2H, d, $J=8.3\text{Hz}$), 8.07 (2H, d, $J=8.3\text{Hz}$)

APCI-MASS : $m/z = 357$ ($\text{M}+\text{H}^+$)

10 Preparation 80

Methyl 4-[4-(6-bromohexyloxy)phenyl]benzoate

IR (KBr) : 2937, 2861, 1724, 1602, 1529, 1436, 1292,
1199, 1112 cm^{-1}

15 NMR (CDCl_3 , δ) : 1.5-2.0 (8H, m), 3.43 (2H, t,
 $J=6.8\text{Hz}$), 3.93 (3H, s), 4.02 (2H, t, $J=6.3\text{Hz}$), 6.98
(2H, d, $J=8.8\text{Hz}$), 7.56 (2H, d, $J=8.8\text{Hz}$), 7.62 (2H,
d, $J=8.4\text{Hz}$), 8.07 (2H, d, $J=8.4\text{Hz}$)

APCI-MASS : $m/z = 391$ ($\text{M}+\text{H}^+$)

20 Preparation 81

4-[4-(5-Bromopentyloxy)phenyl]bromobenzene

IR (KBr) : 2942, 2867, 1604, 1515, 1477, 1286 cm^{-1}

25 NMR (CDCl_3 , δ) : 1.5-2.0 (6H, m), 3.44 (2H, t,
 $J=6.7\text{Hz}$), 3.99 (2H, t, $J=6.2\text{Hz}$), 6.95 (2H, d,
 $J=8.7\text{Hz}$), 7.3-7.6 (6H, m)

APCI-MASS : $m/z = 399$ ($\text{M}+\text{H}^+$)

Preparation 82

30 8-[4-(4-Methoxycarbonylphenyl)phenoxy]octanoyl
piperidine

IR (KBr) : 2935, 2852, 1720, 1639, 1604, 1438,
1292 cm^{-1}

35 NMR (CDCl_3 , δ) : 1.3-1.9 (16H, m), 2.34 (2H, d,
 $J=7.6\text{Hz}$), 3.4-3.6 (4H, m), 3.93 (3H, s), 3.99 (2H,
t, $J=6.4\text{Hz}$), 6.97 (2H, d, $J=8.8\text{Hz}$), 7.55 (2H, d,

J=8.8Hz), 7.61 (2H, d, J=8.6Hz), 8.07 (2H, d, J=8.6Hz)

APCI-MASS : m/z = 438 (M+H⁺)

5 Preparation 83

Methyl 6-[4-(4-n-heptyloxyphenyl)piperazin-1-yl]nicotinate

IR (KBr) : 2933, 2859, 1726, 1608, 1513, 1430, 1280, 1245 cm⁻¹

10 NMR (CDCl₃, δ) : 0.89 (3H, t, J=6.7Hz), 1.2-1.8 (10H, m), 3.17 (4H, t, J=4.9Hz), 3.8-4.0 (9H, m), 6.65 (1H, d, J=9.1Hz), 6.86 (2H, d, J=9.1Hz), 6.96 (2H, d, J=9.1Hz), 8.05 (1H, dd, J=9.1 and 2.3Hz), 8.82 (1H, d, J=2.3Hz)

15 APCI-MASS : m/z = 412 (M+H⁺)

Preparation 84

Methyl 6-[4-[4-(8-bromooctyloxy)phenyl]piperazin-1-yl]nicotinate

20 IR (KBr) : 2933, 2861, 1724, 1608, 1513, 1430, 1280 cm⁻¹

NMR (CDCl₃, δ) : 1.2-2.0 (12H, m), 3.17 (4H, t, J=5.0Hz), 3.40 (2H, t, J=6.8Hz), 3.8-4.0 (9H, m), 6.64 (1H, d, J=9.0Hz), 6.85 (2H, d, J=9.1Hz), 6.96 (2H, d, J=9.1Hz), 8.05 (1H, dd, J=9.0 and 2.2Hz), 8.82 (1H, d, J=2.2Hz)

25 APCI-MASS : m/z = 504 (M+H⁺)

Preparation 85

30 4-[4-(7-Bromoheptyloxy)phenyl]bromobenzene

IR (KBr) : 2935.1, 2856.1, 1604.5 cm⁻¹

NMR (CDCl₃, δ) : 1.18-1.65 (6H, m), 1.70-2.02 (4H, m), 3.41 (2H, t, J=6.8Hz), 3.99 (2H, t, J=6.4Hz), 6.95 (2H, d, J=8.6Hz), 7.40 (2H, d, J=8.6Hz), 7.46 (2H, d, J=8.6Hz), 7.52 (2H, d, J=8.6Hz)

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Preparation 86

4-[4-(8-Bromooctyloxy)phenyl]bromobenzene

NMR (CDCl₃, δ) : 1.22-1.65 (8H, m), 1.65-1.95 (4H, m),
3.41 (2H, t, J=6.8Hz), 3.99 (2H, t, J=6.4Hz), 6.95
(2H, d, J=8.6Hz), 7.40 (2H, d, J=8.6Hz), 7.46 (2H,
d, J=8.6Hz), 7.52 (2H, d, J=8.6Hz)

Preparation 87

Methyl (E)-3-[4-[4-(5-hexenyloxy)phenyl]phenyl]acrylate

NMR (CDCl₃, δ) : 1.50-1.72 (2H, m), 1.72-1.95 (2H, m),
2.05-2.14 (2H, m), 3.82 (3H, s), 4.01 (2H, t,
J=6.3Hz), 4.95-5.10 (2H, m), 5.70-5.93 (1H, m),
6.46 (1H, d, J=16Hz), 6.97 (2H, d, J=8.7Hz), 7.54
(2H, d, J=8.7Hz), 7.58 (4H, s), 7.72 (1H, d,
J=16Hz)

APCI-MASS : m/z = 337 (M⁺+1)

Preparation 88

4-Bromo-4'-(4-methylpentyloxy)biphenyl

IR (KBr) : 2956.3, 2871.5, 1606.4 cm⁻¹

NMR (CDCl₃, δ) : 0.93 (6H, d, J=6.6Hz), 1.25-1.45 (2H,
m), 1.62 (1H, sept, J=6.6Hz), 1.72-1.93 (2H, m),
3.98 (2H, t, J=6.6Hz), 6.95 (2H, d, J=8.6Hz), 7.30-
7.60 (6H, m)

APCI-MASS : m/z = 332, 334 (M⁺, M⁺+2)

The following compounds (Preparations 89 to 90) were
obtained according to a similar manner to that of Preparation
2.

Preparation 89

N-[4-[2-(4-Methylpentyl)-2,3-dihydro-4H-1,2,4-triazol-3-
one-4-yl]phenyl]piperazine ditrifluoroacetate

IR (KBr) : 1668.1, 1519.6, 1203.4, 1176.4, 1130.1 cm⁻¹

NMR (DMSO-d₆, δ) : 0.86 (6H, d, J=6.6Hz), 1.1-1.3 (2H,

m), 1.4-1.8 (3H, m), 3.1-3.3 (4H, m), 3.3-3.5 (4H, m), 3.70 (2H, t, J=7.0Hz), 7.11 (2H, d, J=9.0Hz), 7.53 (2H, d, J=9.0Hz), 8.35 (1H, s), 8.90 (2H, s)

5 Preparation 90

1-(4-Phenylcyclohexyl)piperazine ditrifluoroacetate

IR (KBr) : 1677.8, 1197.6, 1133.9 cm^{-1}

NMR (DMSO- d_6 , δ) : 1.4-1.8 (4H, m), 1.8-2.25 (4H, m),
2.4-2.7 (1H, m), 3.2-3.7 (9H, m), 4.54 (2H, br s),
7.0-7.4 (5H, m), 9.32 (1H, br s)

APCI-MASS : m/z = 245 ($\text{M}^+ + \text{H}$)

The following compounds (Preparations 91 to 103) were
obtained according to a similar manner to that of Preparation
3.

Preparation 91

Methyl 6-[4-(4-octyloxyphenyl)piperazin-1-yl]nicotinate

IR (KBr) : 2923, 1726, 1608, 1515, 1278, 1116 cm^{-1}

NMR (CDCl_3 , δ) : 0.88 (3H, t, J=6.8Hz), 1.2-1.5 (10H, m),
1.7-1.8 (2H, m), 3.1-3.2 (4H, m), 3.8-4.0 (9H, m),
6.64 (1H, d, J=9.0Hz), 6.8-7.0 (4H, m), 8.04 (1H, dd, J=9.0 and 2.4Hz),
8.81 (1H, d, J=2.4Hz)

APCI-MASS : m/z = 426 ($\text{M} + \text{H}^+$)

Preparation 92

4-[4-[4-[2-(4-Methylpentyl)-2,3-dihydro-4H-1,2,4-triazol-3-one-4-yl]phenyl]piperazin-1-yl]benzonitrile

IR (KBr) : 2217.7, 1685.5 cm^{-1}

NMR (CDCl_3 , δ) : 0.90 (6H, d, J=6.6Hz), 1.2-1.4 (2H, m),
1.5-2.0 (3H, m), 3.3-3.4 (4H, m), 3.4-3.6 (4H, m),
3.83 (2H, t, J=7.4Hz), 6.92 (2H, d, J=9.0Hz),
7.01 (2H, d, J=9.0Hz), 7.43 (2H, d, J=9.0Hz), 7.54 (2H, d, J=9.0Hz),
7.62 (1H, s)

Preparation 93

3-Fluoro-4-[4-(4-methoxyphenyl)piperazin-1-yl]benzonitrile

IR (KBr) : 2225.5, 1510.0, 1240.0 cm^{-1}

5 NMR (CDCl_3 , δ) : 3.1-3.55 (8H, m), 3.79 (3H, s),
6.7-7.1 (6H, m), 7.3-7.5 (1H, m)

Preparation 94

10 3-Chloro-4-[4-(4-n-hexyloxyphenyl)piperazin-1-yl]benzonitrile

IR (KBr) : 2223.5, 1592.9, 1510.0, 1490.7, 1236.1 cm^{-1}

15 NMR (CDCl_3 , δ) : 0.90 (3H, t, $J=6.7\text{Hz}$), 1.3-1.6 (6H, m), 1.7-1.9 (2H, m), 3.2-3.4 (8H, m), 3.92 (2H, t, $J=6.6\text{Hz}$), 6.85 (2H, d, $J=9.3\text{Hz}$), 6.94 (2H, d, $J=9.3\text{Hz}$), 7.08 (1H, d, $J=8.4\text{Hz}$), 7.53 (1H, dd, $J=8.4$ and 1.9Hz), 7.64 (1H, d, $J=1.9\text{Hz}$)

APCI-MASS : m/z = 398 ($M^+ + H$)

Preparation 95

20 Ethyl 3-[4-(4-n-hexyloxyphenyl)piperazin-1-yl]-6-pyridazinecarboxylate

IR (KBr) : 1729.8, 1587.1, 1511.9, 1245.8 cm^{-1}

25 NMR (CDCl_3 , δ) : 0.90 (3H, t, $J=6.5\text{Hz}$), 1.2-1.4 (6H, m), 1.44 (3H, t, $J=7.1\text{Hz}$), 1.65-1.85 (2H, m), 3.1-3.25 (4H, m), 3.8-4.0 (6H, m), 4.46 (2H, q, $J=7.1\text{Hz}$), 6.8-7.0 (5H, m), 7.91 (1H, d, $J=9.6\text{Hz}$)

APCI-MASS : m/z = 413 ($M^+ + H$)

Preparation 96

4-(4-Piperidinopiperidin-1-yl)benzonitrile

30 IR (KBr) : 2217.7, 1602.6, 1511.9 cm^{-1}

NMR (CDCl_3 , δ) : 1.35-1.75 (8H, m), 1.92 (2H, d, $J=12.9\text{Hz}$), 2.3-2.6 (5H, m), 2.86 (2H, td, $J=12.8$ and 2.6Hz), 3.90 (2H, d, $J=12.8\text{Hz}$), 6.84 (2H, d, $J=9.1\text{Hz}$), 7.46 (2H, d, $J=9.1\text{Hz}$)

35 APCI-MASS : m/z = 270 ($M^+ + H$)

Preparation 97

5-[4-(4-n-Hexyloxyphenyl)piperazin-1-yl]picolinonitrile

IR (KBr) : 2223.5, 1575.6, 1511.9, 1241.9 cm^{-1}

NMR (CDCl_3 , δ) : 0.90 (3H, t, $J=6.5\text{Hz}$), 1.2-1.55 (6H, m), 1.7-1.85 (2H, m), 3.22 (4H, t, $J=5.1\text{Hz}$), 3.52 (4H, t, $J=5.1\text{Hz}$), 3.92 (2H, t, $J=6.5\text{Hz}$), 6.86 (2H, d, $J=9.4\text{Hz}$), 6.93 (2H, d, $J=9.4\text{Hz}$), 7.13 (1H, dd, $J=8.8$ and 3.0Hz), 7.53 (1H, d, $J=8.8\text{Hz}$), 8.35 (1H, d, $J=3.0\text{Hz}$)

APCI-MASS : $m/z = 365$ ($\text{M}^+ + \text{H}$)

Preparation 98

4-[4-(4-Cyclohexylphenyl)piperazin-1-yl]benzonitrile

IR (KBr) : 2219.7, 1606.4, 1513.8, 1238.1 cm^{-1}

NMR (CDCl_3 , δ) : 1.1-1.5 (6H, m), 1.65-2.0 (4H, m), 2.44 (1H, m), 3.30 (4H, t, $J=5.1\text{Hz}$), 3.46 (4H, t, $J=5.1\text{Hz}$), 6.90 (4H, d, $J=8.9\text{Hz}$), 7.14 (2H, d, $J=8.9\text{Hz}$), 7.52 (2H, d, $J=8.9\text{Hz}$)

APCI-MASS : $m/z = 346$ ($\text{M}^+ + \text{H}$)

Preparation 99

4-[4-(4-n-Hexylphenyl)piperazin-1-yl]benzonitrile

IR (KBr) : 2925.5, 2850.3, 2213.9, 1604.5, 1513.8, 1234.2, 944.9 cm^{-1}

NMR (CDCl_3 , δ) : 0.88 (3H, t, $J=6.4\text{Hz}$), 1.2-1.45 (6H, m), 1.45-1.7 (2H, m), 2.54 (2H, t, $J=7.6\text{Hz}$), 3.2-3.4 (4H, m), 3.4-3.6 (4H, m), 6.89 (2H, d, $J=8.5\text{Hz}$), 6.91 (2H, d, $J=8.9\text{Hz}$), 7.11 (2H, d, $J=8.5\text{Hz}$), 7.52 (2H, d, $J=8.9\text{Hz}$)

Preparation 100

1-[2-(4-n-Hexylphenylamino)ethyl]-2-oxazolidone hydrochloride

IR (KBr) : 2925.5, 2852.2, 1753.0, 1729.8, 1267.0 cm^{-1}

NMR ($\text{DMSO}-d_6$, δ) : 0.85 (3H, t, $J=6.5\text{Hz}$), 1.1-1.4 (6H, m)

m), 1.45-1.7 (2H, m), 2.56 (2H, t, J=7.6Hz), 3.3-3.53 (4H, m), 3.57 (2H, t, J=7.9Hz), 4.24 (2H, t, J=7.9Hz), 7.24 (4H, s)

APCI-MASS : m/z = 291 (M⁺+H)

5

Preparation 101

4-[4-(4-Phenylcyclohexyl)piperazin-1-yl]benzonitrile

IR (KBr) : 2212.0, 1602.6, 1513.8, 1249.6 cm⁻¹

NMR (CDCl₃, δ) : 1.3-1.8 (4H, m), 1.9-2.2 (4H, m),
2.3-2.6 (2H, m), 2.75 (4H, t, J=5.0Hz), 3.34 (4H, t, J=5.0Hz), 6.86 (2H, d, J=8.9Hz), 7.1-7.4 (5H, m), 7.49 (2H, d, J=8.9Hz)

10

APCI-MASS : m/z = 346 (M⁺+H)

15

Preparation 102

Methyl 6-[4-(4-hydroxyphenyl)piperazin-1-yl]nicotinate

IR (KBr) : 3411, 1691, 1602, 1510, 1432, 1249,
1147 cm⁻¹

NMR (DMSO-d₆, δ) : 3.0-3.1 (4H, m), 3.7-3.9 (7H, m),
6.67 (2H, d, J=8.8Hz), 6.84 (2H, d, J=8.8Hz), 6.93 (1H, d, J=9.1Hz), 7.97 (1H, dd, J=2.4 and 9.1Hz), 8.66 (1H, d, J=2.4Hz), 8.88 (1H, s)

20

APCI-MASS : m/z = 314 (M+H)⁺

25

Preparation 103

1-n-Decylindole-5-carboxylic acid

IR (KBr) : 2921, 2854, 1679, 1612, 1427, 1313,
1199 cm⁻¹

NMR (DMSO-d₆, δ) : 0.84 (3H, t, J=6.8Hz), 1.1-1.3 (14H, m), 1.6-1.8 (2H, m), 4.19 (2H, t, J=6.9Hz), 6.57 (1H, s), 7.4-7.8 (3H, m), 8.23 (1H, s), 12.40 (1H, s)

30

APCI-MASS : m/z = 302 (M+H⁺)

35

The following compounds (Preparations 104 to 111) were

obtained according to a similar manner to that of Preparation 10.

Preparation 104

5 (E)-Methyl 4-(4-n-butoxyphenyl)cinnamate
IR (KBr) : 2958, 2939, 2873, 1720, 1637, 1498, 1313,
1195, 1170 cm^{-1}
NMR (CDCl_3 , δ) : 0.98 (3H, t, $J=7.3\text{Hz}$), 1.4-1.8 (4H,
m), 3.81 (3H, s), 4.00 (2H, t, $J=6.4\text{Hz}$), 6.45 (1H,
10 d, $J=16.0\text{Hz}$), 6.97 (2H, d, $J=8.7\text{Hz}$), 7.5-7.7 (6H,
m), 7.72 (1H, d, $J=16.0\text{Hz}$)
APCI-MASS : $m/z = 311$ ($M+H^+$)

Preparation 105

15 Methyl (E)-3-[4-[4-(4-methylpentyloxy)phenyl]phenyl]-
acrylate
IR (KBr) : 2956.3, 2873.4, 1720.2, 1635.3, 1600.6 cm^{-1}
NMR (CDCl_3 , δ) : 0.93 (6H, d, $J=6.5\text{Hz}$), 1.28-1.50 (2H,
m), 1.50-1.95 (3H, m), 3.82 (3H, s), 3.99 (2H, t,
20 $J=6.6\text{Hz}$), 6.44 (1H, d, $J=16.0\text{Hz}$), 6.97 (2H, d,
 $J=8.7\text{Hz}$), 7.49-7.65 (6H, m), 7.71 (1H, d, $J=16\text{Hz}$)
APCI-MASS : $m/z = 339$ (M^++1)

Preparation 106

25 Methyl (E)-3-[4-[4-(6-fluorohexyloxy)phenyl]phenyl]-
acrylate
NMR (CDCl_3 , δ) : 1.23-2.00 (8H, m), 3.81 (3H, s), 4.01
(2H, t, $J=6.4\text{Hz}$), 4.47 (2H, dt, $J=47.4$ and 6.0Hz),
30 6.45 (1H, d, $J=16.0\text{Hz}$), 6.96 (2H, d, $J=8.8\text{Hz}$),
7.45-7.63 (6H, m), 7.72 (1H, d, $J=16.0\text{Hz}$)
APCI-MASS : $m/z = 357$ (M^++1)

Preparation 107

35 Methyl (E)-3-[4-[4-(6-methoxyhexyloxy)phenyl]phenyl]-

acrylate

APCI-MASS : $m/z = 369 (M^+)$

Preparation 108

5 Methyl (E)-3-[4-[4-(8-methoxyoctyloxy)phenyl]phenyl]-
acrylate

IR (KBr) : 2935.1, 2858.0, 1722.1, 1637.3, 1602.6 cm^{-1}

NMR (CDCl_3 , δ) : 1.30-1.70 (10H, m), 1.70-1.92 (2H,
m), 3.33 (3H, s), 3.37 (2H, t, $J=6.5\text{Hz}$), 3.81 (3H,
10 s), 4.00 (2H, t, $J=6.5\text{Hz}$), 6.45 (1H, d, $J=16.0\text{Hz}$),
6.97 (2H, d, $J=8.8\text{Hz}$), 7.46-7.78 (6H, m), 7.72 (1H,
d, $J=16.0\text{Hz}$)

APCI-MASS : $m/z = 397 (M^++1)$

15 Preparation 109

Methyl (E)-3-[4-(4-hydroxyphenyl)phenyl]acrylate

IR (KBr) : 3409.5, 1695.1 cm^{-1}

NMR ($\text{DMSO}-d_6$, δ) : 3.73 (3H, s), 6.64 (1H, d, $J=16\text{Hz}$),
6.85 (2H, d, $J=8.6\text{Hz}$), 7.50-7.83 (5H, m)

20 APCI-MASS : $m/z = 255 (M^++1)$

Preparation 110

Methyl (E)-3-[4-[4-(7-methoxyheptyloxy)phenyl]phenyl]-
acrylate

25 NMR (CDCl_3 , δ) : 1.32-1.70 (8H, m), 1.70-1.92 (2H, m),
3.34 (3H, s), 3.38 (2H, t, $J=6.4\text{Hz}$), 3.81 (3H, s),
4.00 (2H, t, $J=6.5\text{Hz}$), 6.45 (1H, d, $J=16.0\text{Hz}$), 6.97
(2H, d, $J=8.8\text{Hz}$), 7.47-7.65 (6H, m), 7.70 (1H, d,
 $J=16\text{Hz}$)

30 APCI-MASS : $m/z = 383 (M^++1)$

Preparation 111

Methyl (E)-3-[4-[4-(7-fluoroheptyloxy)phenyl]phenyl]-
acrylate

35 IR (KBr) : 2937.1, 2861.8, 1722.1, 1637.3, 1600.6 cm^{-1}

The following compound was obtained according to a similar manner to that of Preparation 20.

Preparation 112

5 Methyl 3-[4-(4-heptylphenyl)phenyl]propanoate

NMR (CDCl₃, δ) : 0.88 (3H, t, J=6.5Hz), 1.15-1.50 (8H, m), 1.50-1.77 (2H, m), 2.52-2.73 (4H, m), 2.99 (2H, t, J=7.8Hz), 3.68 (3H, s), 7.18-7.35 (4H, m), 7.40-7.58 (4H, m)

10 APCI-MASS : m/z = 339 (M⁺+1)

The following compounds (Preparation 113 to 164) were obtained according to a similar manner to that of Preparation 32.

15 Preparation 113

4-(4-Octylphenyl)-2,4-dihydro-3H-1,2,4-triazol-3-one-2-yl-acetic acid

IR (KBr) : 2923.6, 1704.8, 1224.6 cm⁻¹

20 NMR (DMSO-d₆, δ) : 0.85 (3H, t, J=6.7Hz), 1.1-1.4 (10H, m), 1.4-1.7 (2H, m), 2.60 (2H, t, J=7.2Hz), 4.38 (2H, s), 7.32 (2H, d, J=8.5Hz), 7.58 (2H, d, J=8.5Hz), 8.43 (1H, s)

25 Preparation 114

1-Heptyl-4-(4-carboxyphenyl)pyrazole

IR (KBr) : 3106, 2917, 1687, 1612, 1425, 1295, 1184, 952, 860, 773 cm⁻¹

30 NMR (DMSO-d₆, δ) : 0.85 (3H, t, J=6.8Hz), 1.1-1.4 (8H, m), 1.7-1.9 (2H, m), 4.11 (2H, t, J=7.0Hz), 7.69 (2H, d, J=8.5Hz), 7.91 (2H, d, J=8.5Hz), 7.98 (1H, s), 8.32 (1H, s), 12.82 (1H, br)

APCI-MASS : m/z = 287 (M+H⁺)

35 Preparation 115

6-[4-(4-Octyloxyphenyl)piperazin-1-yl]nicotinic acid

IR (KBr pelet) : 2919, 2854, 1697, 1608, 1515, 1429,
1263, 1245, 1228 cm^{-1}

NMR (DMSO- d_6 , δ) : 0.86 (3H, t, $J=6.7\text{Hz}$), 1.1-1.5 (10H,
m), 1.6-1.8 (2H, m), 3.0-3.2 (4H, m), 3.7-3.9 (4H,
m), 3.88 (2H, t, $J=6.4\text{Hz}$), 6.7-7.0 (5H, m), 7.95
(1H, dd, $J=9.0$ and 1.1Hz), 8.64 (1H, d, $J=1.1\text{Hz}$)

APCI-MASS : $m/z = 412$ ($\text{M}+\text{H}^+$)

10 Preparation 116

2-(4-Hexyloxyphenyl)benzoxazole-5-carboxylic acid

IR (KBr) : 2952, 1689, 1677, 1619, 1500, 1415, 1299,
1172, 1024 cm^{-1}

NMR (DMSO- d_6 , δ) : 0.89 (3H, t, $J=6.7\text{Hz}$), 1.2-1.5 (6H,
m), 1.7-1.9 (2H, m), 4.09 (2H, t, $J=6.5\text{Hz}$), 7.16
(2H, d, $J=8.8\text{Hz}$), 7.84 (1H, d, $J=8.5\text{Hz}$), 8.01 (1H,
dd, $J=8.5$ and 1.5Hz), 8.15 (2H, d, $J=8.8\text{Hz}$), 8.26
(1H, d, $J=1.5\text{Hz}$)

APCI-MASS : $m/z = 340$ ($\text{M}+\text{H}^+$)

20 Preparation 117

4-[4-(4-n-Butyloxyphenyl)phenyl]benzoic acid

IR (KBr) : 2958, 2873, 1689, 1600, 1537, 1396 cm^{-1}

25 Preparation 118

6-(4-Heptyloxyphenyl)nicotinic acid

IR (KBr) : 2858, 1699, 1674, 1589, 1425, 1180, 1016,
781 cm^{-1}

NMR (DMSO- d_6 , δ) : 0.87 (3H, t, $J=6.7\text{Hz}$), 1.2-1.5 (8H,
m), 1.6-1.8 (2H, m), 4.04 (2H, t, $J=6.4\text{Hz}$), 7.06
(2H, d, $J=8.9\text{Hz}$), 8.03 (1H, d, $J=8.2\text{Hz}$), 8.13 (2H,
d, $J=8.9\text{Hz}$), 8.27 (1H, dd, $J=8.2$ and 2.2Hz), 9.09
(1H, d, $J=2.2\text{Hz}$), 13.31 (1H, br)

APCI-MASS : $m/z = 314$ ($\text{M}+\text{H}^+$)

Preparation 119

5-(4-Octyloxyphenyl)isoxazole-3-carboxylic acid

IR (KBr pelet) : 2923, 2852, 1704, 1612, 1440, 1272,
1178 cm^{-1}

5 NMR (DMSO- d_6 , δ) : 0.86 (3H, t, $J=6.8\text{Hz}$), 1.2-1.6
(10H, m), 1.6-1.9 (2H, m), 4.03 (2H, t, $J=6.5\text{Hz}$),
7.08 (2H, d, $J=8.9\text{Hz}$), 7.25 (1H, s), 7.86 (2H, d,
 $J=8.9\text{Hz}$)

APCI-MASS : $m/z = 318$ ($\text{M}+\text{H}^+$)

Preparation 120

2-(2-Octyloxypyridin-5-yl)benzoxazole-5-carboxylic acid

IR (KBr) : 2954, 2923, 2854, 1697, 1683, 1625, 1488,
1290 cm^{-1}

15 NMR (DMSO- d_6 , δ) : 0.86 (3H, t, $J=7.6\text{Hz}$), 1.2-1.5
(10H, m), 1.7-1.8 (2H, m), 4.36 (2H, t, $J=6.6\text{Hz}$),
7.04 (1H, d, $J=8.7\text{Hz}$), 7.88 (1H, d, $J=8.5\text{Hz}$), 8.04
(1H, dd, $J=8.5$ and 1.6Hz), 8.29 (1H, d, $J=1.6\text{Hz}$),
8.43 (1H, dd, $J=8.7$ and 2.4Hz), 8.99 (1H, d,
20 $J=2.4\text{Hz}$), 13.0-13.2 (1H, br)

APCI-MASS : $m/z = 369$ ($\text{M}+\text{H}^+$)

Preparation 121

2-[4-(4-Hexylphenyl)phenyl]benzoxazole-5-carboxylic acid

25 IR (KBr) : 2923, 2854, 1683, 1411, 1299, 1054 cm^{-1}

APCI-MASS : $m/z = 400$ ($\text{M}+\text{H}^+$)

Preparation 122

6-[4-(4-n-Butyloxyphenyl)phenyl]nicotinic acid

30 IR (KBr) : 3406, 2958, 1691, 1591, 1394, 1284,
1253 cm^{-1}

NMR (DMSO- d_6 , δ) : 0.94 (3H, t, $J=7.3\text{Hz}$), 1.4-1.8 (4H,
m), 4.01 (2H, t, $J=6.4\text{Hz}$), 7.02 (2H, d, $J=8.7\text{Hz}$),
7.57 (2H, d, $J=8.7\text{Hz}$), 7.61 (2H, d, $J=8.2\text{Hz}$), 7.83
35 (2H, d, $J=8.2\text{Hz}$), 8.05 (1H, d, $J=8.5\text{Hz}$), 8.22 (1H,

dd, J=8.5 and 1.6Hz), 9.14 (1H, d, J=1.6Hz)
APCI-MASS : m/z = 348 (M+H⁺)

Preparation 123

5 4-[4-(5-Phenoxypropyloxy)phenyl]benzoic acid
NMR (DMSO-d₆, δ) : 1.5-1.7 (2H, m), 1.7-1.9 (4H, m),
3.98 (2H, t, J=6.3Hz), 4.05 (2H, t, J=6.1Hz), 6.8-
7.0 (3H, m), 7.05 (2H, d, J=8.6Hz), 7.25 (2H, t,
J=8.2Hz), 7.68 (2H, d, J=8.5Hz), 7.75 (2H, d,
10 J=8.2Hz), 7.98 (2H, d, J=8.2Hz), 12.8-13.0 (1H, br
s)
APCI-MASS : m/z = 375 (M-H)⁻

Preparation 124

15 4-[5-(4-n-Hexyloxyphenyl)-1,3,4-oxadiazol-2-yl]benzoic
acid
IR (KBr) : 2935, 2854, 1685, 1612, 1495, 1425, 1286,
1251 cm⁻¹
NMR (DMSO-d₆, δ) : 0.89 (3H, t, J=6.7Hz), 1.2-1.5 (6H,
20 m), 1.6-1.9 (3H, m), 4.12 (2H, t, J=6.4Hz), 7.19
(2H, d, J=8.7Hz), 8.08 (2H, d, J=8.7Hz), 8.18 (2H,
d, J=8.4Hz), 8.24 (2H, d, J=8.4Hz)
APCI-MASS : m/z = 367 (M+H)⁺

Preparation 125

25 4-[5-(4-n-Hexyloxyphenyl)-1,3,4-thiadiazol-2-yl]benzoic
acid
IR (KBr) : 2952, 2586, 1699, 1604, 1517, 1432, 1251,
1174 cm⁻¹
30 NMR (DMSO-d₆, δ) : 0.89 (3H, t, J=6.7Hz), 1.3-1.9 (8H,
m), 4.04 (2H, t, J=6.3Hz), 7.13 (2H, d, J=8.8Hz),
7.97 (2H, d, J=8.8Hz), 8.11 (4H, s)
APCI-MASS : m/z = 383 (M+H)⁺

35 Preparation 126

5-(4-Octyloxyphenyl)-1-methylpyrazole-3-carboxylic acid

IR (KBr pelet) : 2950, 2923, 1695, 1450, 1282, 1251,
956 cm^{-1}

NMR (DMSO-d_6 , δ) : 0.86 (3H, t, $J=6.8\text{Hz}$), 1.2-1.5
(10H, m), 1.6-1.8 (2H, m), 3.98 (2H, t, $J=6.5\text{Hz}$),
4.10 (3H, s), 6.95 (1H, d, $J=8.8\text{Hz}$), 7.18 (1H, s),
7.73 (2H, d, $J=8.8\text{Hz}$), 13.37 (1H, br)

APCI-MASS : $m/z = 331$ ($\text{M}+\text{H}^+$)

10 Preparation 127

4-[3-(4-n-Pentyloxyphenyl)pyrazol-5-yl]benzoic acid

IR (KBr) : 3224, 2956, 1692, 1614, 1506, 1251 cm^{-1}

NMR (DMSO-d_6 , δ) : 0.91 (3H, t, $J=6.9\text{Hz}$), 1.3-1.5 (4H,
m), 1.6-1.8 (2H, m), 4.00 (2H, t, $J=6.5\text{Hz}$), 7.02
(2H, d, $J=8.8\text{Hz}$), 7.19 (1H, s), 7.75 (2H, d,
 $J=8.8\text{Hz}$), 7.95 (2H, d, $J=8.7\text{Hz}$), 8.02 (2H, d,
 $J=8.7\text{Hz}$), 12.8-13.3 (2H, br)

APCI-MASS : $m/z = 351$ ($\text{M}+\text{H}^+$)

20 Preparation 128

5-[4-(4-n-Butoxyphenyl)phenyl]furan-2-carboxylic acid

IR (KBr) : 2958, 2873, 1679, 1487, 1253, 1166 cm^{-1}

NMR (DMSO-d_6 , δ) : 0.95 (3H, t, $J=7.3\text{Hz}$), 1.3-1.8 (4H,
m), 4.02 (2H, t, $J=6.3\text{Hz}$), 7.03 (2H, d, $J=8.6\text{Hz}$),
7.17 (1H, d, $J=3.6\text{Hz}$), 7.33 (1H, d, $J=3.6\text{Hz}$), 7.66
(2H, d, $J=8.6\text{Hz}$), 7.74 (2H, d, $J=8.4\text{Hz}$), 7.86 (2H,
d, $J=8.4\text{Hz}$), 13.1 (1H, br s)

APCI-MASS : $m/z = 337$ ($\text{M}+\text{H}^+$)

30 Preparation 129

3-(S)-Hydroxyhexadecanoic acid

IR (KBr) : 1679.7, 1467.6, 1224.6 cm^{-1}

NMR (CDCl_3 , δ) : 0.88 (3H, t, $J=6.4\text{Hz}$), 1.1-1.7 (24H,
m), 2.35-2.65 (2H, m), 4.03 (1H, m), 5.41 (1H, br
s)

Preparation 130

6-[4-(4-n-Hexyloxyphenyl)piperazin-1-yl]pyridazine-3-carboxylic acid

IR (KBr) : 1697.1, 1589.1, 1515.8, 1448.3 cm^{-1}

5 NMR (DMSO- d_6 , δ) : 0.87 (3H, t, $J=6.4\text{Hz}$), 1.2-1.5 (6H, m), 1.6-1.8 (2H, m), 3.0-3.2 (4H, m), 3.7-4.0 (6H, m), 6.83 (2H, d, $J=9.0\text{Hz}$), 6.95 (2H, d, $J=9.0\text{Hz}$), 7.36 (1H, d, $J=9.6\text{Hz}$), 7.86 (1H, d, $J=9.6\text{Hz}$), 11.68 (1H, s)

10

Preparation 131

4-[4-[1-(4-n-Hexyloxyphenyl)piperidin-4-yl]piperazin-1-yl]benzoic acid hydrochloride

IR (KBr) : 1699.0, 1608.3, 1513.8 cm^{-1}

15 NMR (DMSO- d_6 , δ) : 0.88 (3H, t, $J=6.5\text{Hz}$), 1.2-1.5 (6H, m), 1.6-1.8 (2H, m), 2.0-2.45 (3H, m), 3.2-3.8 (12H, m), 3.94 (2H, t, $J=6.4\text{Hz}$), 4.03 (2H, d, $J=11\text{Hz}$), 6.95 (2H, d, $J=8.7\text{Hz}$), 7.07 (2H, d, $J=8.9\text{Hz}$), 7.32 (2H, br s), 7.83 (2H, d, $J=8.9\text{Hz}$)

20 APCI-MASS : $m/z = 466$ ($\text{M}^+ + \text{H}$)

Preparation 132

6-(8-Methoxyoctyloxy)-2-naphthoic acid

IR (KBr) : 2937.1, 2854.1, 1677.8, 1211.1 cm^{-1}

25 NMR (DMSO- d_6 , δ) : 1.2-1.6 (10H, m), 1.7-1.9 (2H, m), 3.20 (3H, s), 3.29 (2H, t, $J=6.5\text{Hz}$), 4.11 (2H, t, $J=6.4\text{Hz}$), 7.23 (1H, dd, $J=9.0$ and 2.3Hz), 7.39 (1H, d, $J=2.3\text{Hz}$), 7.85 (1H, d, $J=8.7\text{Hz}$), 7.93 (1H, d, $J=8.7\text{Hz}$), 7.99 (1H, d, $J=9.0\text{Hz}$), 8.51 (1H, s), 12.9 (1H, s)

30

Preparation 133

Mixture of (E) and (Z)-3-[4-(4-Heptylphenyl)phenyl]-2-butenic acid

35 NMR (CDCl_3 , δ) : 0.88 (3H, t, $J=6.6\text{Hz}$), 1.15-1.50 (8H,

m), 1.52-1.75 (2H, m), 2.63 and 3.62 (total 3H, each s), 2.53-2.75 (2H, m), 6.24 and 5.68 (total 1H, each s), 7.19-7.35 (2H, m), 7.47-7.70 (6H, m)
APCI-MASS : $m/z = 337 (M^+ + 1)$, 351 (methyl ester⁺+1)

5

Preparation 134

3-[4-(4-Heptylphenyl)phenyl]propanoic acid

NMR (CDCl₃, δ) : 0.88 (3H, t, J=6.6Hz), 1.13-1.48 (8H, m), 1.48-1.75 (2H, m), 2.52-2.83 (4H, m), 3.00 (2H, t, J=7.8Hz), 7.15-7.35 (4H, m), 7.40-7.60 (4H, m)

10

APCI-MASS : $m/z = 323 (M^+ - 1)$

Preparation 135

4-(4-n-Heptylphenyl)benzoyl-carboxylic acid

NMR (CDCl₃, δ) : 0.88 (3H, t, J=6.6Hz), 1.13-1.50 (8H, m), 1.50-1.75 (2H, m), 2.66 (2H, t, J=7.7Hz), 7.20-7.40 (2H, m), 7.50-7.66 (2H, m), 7.66-7.84 (2H, m), 8.40-8.60 (2H, m)

15

APCI-MASS : $m/z = 323 (M^+ - 1)$

20

Preparation 136

6-Hexylnaphthalene-2-carboxylic acid

NMR (CDCl₃, δ) : 0.89 (3H, t, J=6.8Hz), 1.15-1.53 (6H, m), 1.55-1.84 (2H, m), 2.80 (2H, t, J=7.6Hz), 7.42 (1H, dd, J=1.7 and 8.4Hz), 7.67 (1H, s), 7.84 (1H, d, J=8.6Hz), 7.90 (1H, d, J=8.4Hz), 8.09 (1H, dd, J=1.7 and 8.6Hz), 8.68 (1H, s)

25

APCI-MASS : $m/z = 257 (M^+ + 1)$, 271 (methyl ester⁺+1)

30 Preparation 137

3-(E)-[4-[4-(7-Methoxyheptyloxy)phenyl]phenyl]acrylic acid

NMR (DMSO-d₆, δ) : 1.20-1.60 (8H, m), 1.60-1.83 (2H, m), 3.21 (3H, s), 3.25-3.60 (2H, m), 4.01 (2H, t, J=6.4Hz), 6.54 (1H, d, J=16.0Hz), 7.02 (2H, d,

35

J=8.8Hz), 7.55-7.80 (7H, m)

APCI-MASS : $m/z = 369 (M^+ + 1)$

Preparation 138

5 3-(E)-[4-[4-(8-Methoxyoctyloxy)phenyl]phenyl]acrylic
acid

IR (KBr) : 3037.3, 2933.2, 2858.0, 2551.4, 1706.7,
1677.8, 1629.6, 1602.6 cm^{-1}

NMR (DMSO- d_6 , δ) : 1.18-1.55 (10H, m), 1.65-1.83 (2H,
10 m), 3.18-3.45 (5H, m), 4.01 (2H, t, J=6.5Hz), 6.53
(1H, d, J=16.0Hz), 7.02 (2H, d, J=8.8Hz), 7.50-8.80
(7H, m)

APCI-MASS : $m/z = 383 (M^+ + 1)$

15 Preparation 139

3-(E)-[4-[4-(5-Hexenyloxy)phenyl]phenyl]acrylic acid

NMR (DMSO- d_6 , δ) : 1.42-1.63 (2H, m), 1.63-1.85 (2H,
m), 2.00-2.20 (2H, m), 4.03 (2H, t, J=6.3Hz), 4.90-
5.15 (2H, m), 5.68-5.97 (1H, m), 6.54 (1H, d,
20 J=16Hz), 7.02 (2H, d, J=8.7Hz), 7.50-7.80 (7H, m)

APCI-MASS : $m/z = 323 (M^+ + 1)$

Preparation 140

25 3-(E)-[4-[4-(4-Methylpentyloxy)phenyl]phenyl]acrylic
acid

IR (KBr) : 2956.3, 2869.6, 2713.4, 2599.6, 1689.3,
1627.6, 1602.6 cm^{-1}

NMR (DMSO- d_6 , δ) : 0.89 (6H, d, J=6.5Hz), 1.15-1.43
(2H, m), 1.48-1.90 (3H, m), 4.00 (2H, t, J=6.7Hz),
30 6.54 (1H, d, J=16Hz), 7.02 (2H, d, J=8.7Hz), 7.50-
7.90 (7H, m)

APCI-MASS : $m/z = 325 (M^+ + 1)$

Preparation 141

35 3-(E)-[4-[4-(6-Fluorohexyloxy)phenyl]phenyl]acrylic acid

NMR (CDCl₃, δ) : 1.39-2.00 (8H, m), 4.01 (2H, t, J=6.5Hz), 4.47 (2H, dt, J=47.3 and 6.0Hz), 6.49 (1H, d, J=15.9Hz), 6.98 (2H, d, J=8.7Hz), 7.40-7.70 (6H, m), 7.81 (1H, d, J=15.9Hz)

5 APCI-MASS : m/z = 343 (M⁺+1)

Preparation 142

3-(E)-[4-[4-(6-Methoxyhexyloxy)phenyl]phenyl]acrylic acid

10 NMR (DMSO-d₆, δ) : 1.22-1.63 (6H, m), 1.63-1.88 (2H, m), 3.21 (3H, s), 3.22-3.40 (2H, m), 4.00 (2H, t, J=6.5Hz), 6.54 (1H, d, J=15.8Hz), 7.02 (2H, d, J=8.7Hz), 7.50-7.84 (7H, m)

APCI-MASS : m/z = 369 (methyl ester, M⁺+1)

15

Preparation 143

4-[4-[8-(Tetrahydropyran-2-yl-oxy)octyloxy]phenyl]benzoic acid

20 IR (KBr) : 2935, 1697, 1683, 1604, 1303, 1290, 1197 cm⁻¹

NMR (DMSO-d₆, δ) : 1.2-1.8 (18H, m), 3.3-3.9 (4H, m), 4.01 (2H, t, J=6.3Hz), 4.5-4.6 (1H, m), 7.03 (2H, d, J=8.7Hz), 7.67 (2H, d, J=8.7Hz), 7.74 (2H, d, J=8.3Hz), 7.98 (2H, d, J=8.3Hz)

25 APCI-MASS : m/z = 425 (M-H⁺)

Preparation 144

4-[3-(4-n-Hexyloxyphenyl)pyrazol-5-yl]benzoic acid

30 IR (KBr) : 2956, 2935, 1693, 1614, 1508, 1432, 1251, 1178 cm⁻¹

NMR (DMSO-d₆, δ) : 0.89 (3H, t, J=6.4Hz), 1.2-1.5 (6H, m), 1.6-1.8 (2H, m), 4.00 (2H, t, J=6.4Hz), 7.02 (2H, d, J=8.7Hz), 7.12 (1H, s), 7.74 (2H, d, J=8.7Hz), 7.95 (2H, d, J=8.8Hz), 8.01 (2H, d, J=8.8Hz), 13.17 (1H, s)

35

APCI-MASS : $m/z = 365$ ($M+H^+$)

Preparation 145

4-[4-[4-(6-Methoxyhexyloxy)phenyl]phenyl]benzoic acid

5 IR (KBr) : 2939, 2861, 1685, 1602, 1430, 1286,
1128 cm^{-1}

NMR (DMSO-d_6 , δ) : 1.3-1.8 (8H, m), 3.21 (3H, s),
3.3-3.4 (2H, m), 4.01 (2H, t, $J=6.5\text{Hz}$), 7.04 (2H,
d, $J=8.6\text{Hz}$), 7.66 (2H, d, $J=8.6\text{Hz}$), 7.7-7.9 (6H,
10 m), 8.03 (2H, d, $J=8.2\text{Hz}$)

APCI-MASS : $m/z = 405$ ($M+H^+$)

Preparation 146

4-[5-[4-(8-Methoxyoctyloxy)phenyl]-1,3,4-thiadiazol-2-
15 yl]benzoic acid

IR (KBr) : 2931, 2854, 1691, 1602, 1251 cm^{-1}

NMR (DMSO-d_6 , δ) : 1.2-2.0 (12H, m), 3.20 (3H, s),
3.29 (2H, t, $J=6.4\text{Hz}$), 4.04 (2H, t, $J=6.4\text{Hz}$), 7.13
(2H, t, $J=8.8\text{Hz}$), 7.9-8.2 (6H, m), 13.95 (1H, br)

20 APCI-MASS : $m/z = 441$ ($M+H^+$)

Preparation 147

4-(4-n-Butoxyphenyl)cinnamic acid

IR (KBr) : 2958, 2871, 1695, 1625, 1498, 1249 cm^{-1}

25 NMR (DMSO-d_6 , δ) : 0.94 (3H, t, $J=7.3\text{Hz}$), 1.44 (2H,
tq, $J=7.0$ and 7.3Hz), 1.71 (2H, tt, $J=7.0$ and
6.4Hz), 4.01 (2H, t, $J=6.4\text{Hz}$), 6.54 (1H, d,
 $J=16.0\text{Hz}$), 7.02 (2H, d, $J=8.7\text{Hz}$), 7.6-7.9 (7H, m)

APCI-MASS : $m/z = 297$ ($M+H^+$)

30

Preparation 148

4-[5-(4-Cyclohexylphenyl)-1,3,4-thiadiazol-2-yl]benzoic
acid

IR (KBr) : 2925, 2850, 1683, 1429, 1292 cm^{-1}

35 NMR (DMSO-d_6 , δ) : 1.1-1.5 (5H, m), 1.6-2.0 (5H, m),

APCI-MASS : $m/z = 365 \text{ (M+H)}^+$

4-[5-[4-(Piperidin-1-yl)phenyl]-1,3,4-thiadiazol-2-yl]-
benzoic acid

NMR (DMSO- d_6 , δ) : 1.61 (6H, s), 3.31 (4H, s), 7.05

APCI-MASS : $m/z = 366 \text{ (M+H)}^+$

15 4-[5-[4-[4-n-Propyloxyphenyl)phenyl]-1,3,4-oxadiazol-2-
yl]benzoic acid

NMR (DMSO- d_6 , δ) : 1.00 (3H, t, $J=7.3\text{Hz}$), 1.76 (2H,

APCI-MASS : $m/z = 401 \text{ (M+H)}^+$

25 4-(5-n-Nonyl-1,3,4-oxadiazol-2-yl)benzoic acid

NMR (DMSO- d_6 , δ) : 0.84 (3H, t, $J=6.5\text{Hz}$), 1.2-1.5

APCI-MASS : $m/z = 317$ ($M+H^+$)

4-[3-(4-n-Hexyloxyphenyl)-1,2,4-oxadiazol-5-yl]benzoic
acid

35 IR (KBr) : 2942, 2869, 1695, 1421, 1251 cm^{-1}

NMR (DMSO- d_6 , δ) : 0.89 (3H, t, $J=6.8\text{Hz}$), 1.2-1.8 (8H, m), 4.06 (2H, t, $J=6.5\text{Hz}$), 7.13 (2H, d, $J=8.9\text{Hz}$), 8.03 (2H, d, $J=8.9\text{Hz}$), 8.17 (2H, d, $J=8.5\text{Hz}$), 8.28 (2H, d, $J=8.5\text{Hz}$)

5 APCI-MASS : $m/z = 367$ ($M+H$)⁺

Preparation 153

4-[4-[4-(5-Methoxypentyloxy)phenyl]phenyl]phenylacetic acid

10 IR (KBr) : 2939, 2861, 1699, 1253, 1182, 1124 cm^{-1}

NMR (DMSO- d_6 , δ) : 1.4-1.9 (6H, m), 3.22 (3H, s), 3.39 (2H, t, $J=6.2\text{Hz}$), 3.61 (2H, s), 4.01 (2H, t, $J=6.4\text{Hz}$), 7.02 (2H, d, $J=8.8\text{Hz}$), 7.35 (2H, d, $J=8.2\text{Hz}$), 7.6-7.8 (8H, m)

15 APCI-MASS : $m/z = 405$ ($M+H$)⁺

Preparation 154

4-[5-(4-n-Octyloxyphenyl)-1,3,4-thiadiazol-2-yl]benzoic acid

20 IR (KBr) : 2921, 2856, 1691, 1432, 1251 cm^{-1}

NMR (DMSO- d_6 , δ) : 0.87 (3H, t, $J=6.7\text{Hz}$), 1.2-1.5 (10H, m), 1.7-1.9 (2H, m), 4.07 (2H, t, $J=6.5\text{Hz}$), 7.13 (2H, d, $J=8.9\text{Hz}$), 7.97 (2H, d, $J=8.9\text{Hz}$), 8.12 (4H, s)

25 APCI-MASS : $m/z = 411$ ($M+H$)⁺

Preparation 155

4-[5-(4-Trans-n-pentylcyclohexyl)-1,3,4-thiadiazol-2-yl]benzoic acid

30 IR (KBr) : 2919, 2848, 1677, 1430, 1294 cm^{-1}

NMR (DMSO- d_6 , δ) : 0.87 (3H, t, $J=6.9\text{Hz}$), 1.0-1.4 (11H, m), 1.5-1.6 (2H, m), 1.8-2.0 (2H, m), 2.1-2.3 (2H, m), 3.1-3.3 (1H, m), 8.07 (4H, s)

APCI-MASS : $m/z = 359$ ($M+H$)⁺

Preparation 156

4-[3-(4-n-Pentyloxyphenyl)isoxazol-5-yl]benzoic acid

IR (KBr) : 2925, 2869, 1699, 1687, 1612, 1432, 1251,
1178 cm^{-1}

5 NMR (DMSO-d_6 , δ) : 0.91 (3H, t, $J=6.9\text{Hz}$), 1.2-1.5 (4H,
m), 1.7-1.9 (2H, m), 4.04 (2H, t, $J=6.5\text{Hz}$), 7.09
(2H, d, $J=8.8\text{Hz}$), 7.69 (1H, s), 7.85 (2H, d,
 $J=8.8\text{Hz}$), 8.01 (2H, d, $J=8.5\text{Hz}$), 8.11 (2H, d,
 $J=8.5\text{Hz}$)

10 APCI-MASS : $m/z = 352$ ($\text{M}+\text{H}^+$)

Preparation 157

4-[5-[4-(8-Methoxyoctyloxy)phenyl]-1,3,4-oxadiazol-2-
yl]benzoic acid

15 IR (KBr) : 2967, 2937, 2877, 1687, 1290 cm^{-1}
NMR (DMSO-d_6 , δ) : 1.2-1.6 (10H, m), 1.7-1.9 (2H, m),
3.20 (3H, s), 3.29 (2H, t, $J=6.4\text{Hz}$), 4.08 (2H, t,
 $J=6.5\text{Hz}$), 7.17 (2H, d, $J=8.9\text{Hz}$), 8.07 (2H, d,
 $J=8.9\text{Hz}$), 8.15 (2H, d, $J=8.6\text{Hz}$), 8.24 (2H, d,
20 $J=8.6\text{Hz}$)

APCI-MASS : $m/z = 425$ ($\text{M}+\text{H}$)⁺

Preparation 158

4-[4-(6-Phenylpyridazin-3-yl-oxy)phenyl]benzoic acid

25 IR (KBr) : 1700, 1687, 1608, 1427, 1284, 1186 cm^{-1}
NMR (DMSO-d_6 , δ) : 7.40 (2H, d, $J=8.6\text{Hz}$), 7.5-7.7 (4H,
m), 7.7-7.9 (4H, m), 7.9-8.1 (4H, m), 8.35 (1H, d,
 $J=9.2\text{Hz}$), 12.99 (1H, br s)

APCI-MASS : $m/z = 369$ ($\text{M}+\text{H}$)⁺

30

Preparation 159

4-[5-(4-n-Octyloxyphenyl)-1,3,4-oxadiazol-2-yl]benzoic
acid

35 IR (KBr) : 2921, 2852, 1685, 1612, 1496, 1425, 1288,
1251 cm^{-1}

080973 0519
"61250" 226080

NMR (DMSO-d₆, δ) : 0.87 (3H, t, J=6.7Hz), 1.2-1.5
(10H, m), 1.7-1.9 (2H, m), 4.08 (2H, t, J=6.4Hz),
7.17 (2H, d, J=8.7Hz), 8.07 (2H, d, J=8.7Hz), 8.15
(2H, d, J=8.5Hz), 8.24 (2H, d, J=8.5Hz), 13.36 (1H,
br)
APCI-MASS : m/z = 395 (M+H⁺)

Preparation 160

4-[2-(4-n-Hexyloxyphenyl)pyrimidin-6-yl]benzoic acid

IR (KBr) : 2944, 2863, 1697, 1585, 1415, 1386,
1253 cm⁻¹

NMR (DMSO-d₆, δ) : 0.89 (3H, t, J=6.7Hz), 1.2-1.6 (6H,
m), 1.7-1.9 (2H, m), 4.07 (2H, t, J=6.6Hz), 7.10
(2H, d, J=8.9Hz), 8.00 (1H, d, J=5.2Hz), 8.13 (2H,
d, J=8.4Hz), 8.44 (2H, d, J=5.9Hz), 8.47 (2H, d,
J=5.9Hz), 8.95 (1H, d, J=5.2Hz)

APCI-MASS : m/z = 377 (M+H⁺)

Preparation 161

4-[4-(7-Piperidinocarbonylheptyloxy)phenyl]benzoic acid

IR (KBr) : 2933, 2858, 1697, 1677, 1637, 1604, 1429,
1249 cm⁻¹

NMR (DMSO-d₆, δ) : 1.2-1.8 (16H, m), 2.26 (2H, t,
J=7.5Hz), 3.2-3.5 (4H, m), 4.01 (2H, t, J=6.4Hz),
7.03 (2H, d, J=8.8Hz), 7.67 (2H, d, J=8.8Hz), 7.74
(2H, d, J=8.4Hz), 7.98 (2H, d, J=8.4Hz)

APCI-MASS : m/z = 424 (M+H⁺)

Preparation 162

6-[4-(4-n-Heptyloxyphenyl)piperazin-1-yl]nicotinic acid

IR (KBr) : 2929, 2854, 1695, 1673, 1606, 1577, 1515,
1421, 1245 cm⁻¹

NMR (DMSO-d₆, δ) : 0.86 (3H, t, J=6.7Hz), 1.2-1.5 (8H,
m), 1.6-1.8 (2H, m), 3.0-3.2 (4H, m), 3.6-3.8 (4H,
m), 3.87 (2H, t, J=6.5Hz), 6.8-7.2 (5H, m), 7.95

(1H, dd, J=8.9 and 2.3Hz), 8.62 (1H, d, J=2.3Hz)

APCI-MASS : m/z = 398 (M+H⁺)

Preparation 163

5 6-[4-[4-(8-Methoxyoctyloxy)phenyl]piperazin-1-yl]-
nicotinic acid

IR (KBr) : 2933, 2856, 1697, 1672, 1605, 1511, 1421,
1245 cm⁻¹

10 NMR (DMSO-d₆, δ) : 1.2-1.8 (12H, m), 3.08 (4H, t,
J=5.0Hz), 3.20 (3H, s), 3.28 (2H, t, J=6.5Hz), 3.78
(4H, t, J=4.6Hz), 3.87 (2H, t, J=6.4Hz), 6.8-7.0
(5H, m), 7.95 (1H, dd, J=9.0 and 2.2Hz), 8.65 (1H,
d, J=2.2Hz), 12.54 (1H, s)

APCI-MASS : m/z = 442 (M+H⁺)

15

Preparation 164

4-[5-[4-(4-n-Propyloxyphenyl)phenyl]-1,3,4-thiadiazol-2-
yl]benzoic acid

IR (KBr) : 1685, 1537, 1423, 817 cm⁻¹

20 NMR (DMSO-d₆, δ) : 1.00 (3H, t, J=6.7Hz), 1.6-1.8 (2H,
m), 4.00 (2H, t, J=6.6Hz), 7.0-7.2 (2H, d,
J=8.6Hz), 7.6-8.1 (10H, m)

APCI-MASS : m/z = 417 (M+H)⁺

25 Preparation 165

To a solution of Ethyl 4-[5-(4-n-pentyloxyphenyl)-
isoxazol-3-yl]benzoate (6.33 g) in ethanol (60 ml) and
tetrahydrofuran (90 ml) was added 2N sodium hydroxide aqueous
solution (12.5 ml) at 80°C. The mixture was refluxed for 1
30 hour and poured into ice-water. The suspension was adjusted
to pH 2.0 with 1N HCl. The precipitate was collected by
filtration, washed with water and dried to give 4-[5-(4-n-
pentyloxyphenyl)isoxazol-3-yl]benzoic acid (5.80 g).

35 IR (KBr) : 2939, 2867, 1681, 1614, 1429, 1255, 1178,
821 cm⁻¹

00009723 05219
267250 2260880

NMR (DMSO- d_6 , δ) : 0.91 (3H, t, $J=7.1\text{Hz}$), 1.3-1.5 (4H, m), 1.6-1.8 (2H, m), 4.04 (2H, t, $J=6.5\text{Hz}$), 7.11 (2H, d, $J=8.9\text{Hz}$), 7.54 (1H, s), 7.85 (2H, d, $J=8.9\text{Hz}$), 7.98 (2H, d, $J=8.6\text{Hz}$), 8.11 (2H, d, $J=8.6\text{Hz}$)

APCI-MASS : $m/z = 352$ ($M+H$)⁺

The following compounds (Preparations 166 to 170) were obtained according to a similar manner to that of Preparation 40.

Preparation 166

5-[4-(4-n-Hexyloxyphenyl)piperazin-1-yl]picolic acid trihydrochloride

IR (KBr) : 1689.3, 1577.5, 1511.9, 1241.9 cm^{-1}

NMR (DMSO- d_6 , δ) : 0.88 (3H, t, $J=6.5\text{Hz}$), 1.15-1.5 (6H, m), 1.6-1.8 (2H, m), 3.1-3.25 (4H, m), 3.45-3.6 (4H, m), 3.89 (2H, t, $J=6.4\text{Hz}$), 6.84 (2H, d, $J=9.1\text{Hz}$), 6.97 (2H, d, $J=9.1\text{Hz}$), 7.43 (1H, dd, $J=8.8$ and 3.0Hz), 7.90 (1H, dd, $J=8.8$ and 0.7Hz), 8.41 (1H, dd, $J=3.0$ and 0.7Hz)

APCI-MASS : $m/z = 384$ ($M^+ + H$)

Preparation 167

4-[4-(4-Phenylcyclohexyl)piperazin-1-yl]benzoic acid dihydrochloride

IR (KBr) : 1700.9, 1606.4, 1220.7, 1180.2 cm^{-1}

NMR (DMSO- d_6 , δ) : 1.4-1.85 (4H, m), 1.9-2.05 (2H, m), 2.2-2.4 (2H, m), 3.1-3.5 (6H, m), 3.5-3.7 (2H, m), 3.9-4.2 (2H, m), 7.06 (2H, d, $J=8.8\text{Hz}$), 7.1-7.4 (5H, m), 7.83 (2H, d, $J=8.8\text{Hz}$)

APCI-MASS : $m/z = 365$ ($M^+ + H$)

Preparation 168

4-(4-Trans-n-pentylcyclohexyl)benzoic acid

IR (KBr) : 1681.6, 1423.2, 1290.1 cm^{-1}

NMR (CDCl_3 , δ) : 0.90 (3H, t, $J=6.6\text{Hz}$), 1.0-1.6 (13H, m), 1.89 (4H, d, $J=10\text{Hz}$), 2.54 (1H, t, $J=12\text{Hz}$), 7.30 (2H, d, $J=8.3\text{Hz}$), 8.03 (2H, d, $J=8.3\text{Hz}$)

5 APCI-MASS : $m/z = 274$ ($M^+ + H$)

Preparation 169

4-(4-Piperidinopiperidin-1-yl)benzoic acid

IR (KBr) : 1710.6, 1403.9 cm^{-1}

10 NMR ($\text{DMSO}-d_6$, δ) : 1.6-2.1 (8H, m), 2.17 (2H, d, $J=12\text{Hz}$), 2.7-3.05 (4H, m), 3.2-3.5 (1H, m), 3.35 (2H, d, $J=12\text{Hz}$), 4.05 (2H, d, $J=13\text{Hz}$), 7.01 (2H, d, $J=8.9\text{Hz}$), 7.77 (2H, d, $J=8.9\text{Hz}$), 10.84 (1H, s)

APCI-MASS : $m/z = 289$ ($M^+ + H$)

15

Preparation 170

3-Chloro-4-[4-(4-n-hexyloxyphenyl)piperazin-1-yl]benzoic acid dihydrochloride

IR (KBr) : 1712.5, 1598.7, 1513.8, 1251.6 cm^{-1}

20 NMR ($\text{DMSO}-d_6$, δ) : 0.88 (3H, t, $J=6.6\text{Hz}$), 1.2-1.5 (6H, m), 1.6-1.8 (2H, m), 3.4-3.6 (8H, m), 3.98 (2H, t, $J=6.4\text{Hz}$), 7.02 (2H, d, $J=9.0\text{Hz}$), 7.32 (1H, d, $J=8.1\text{Hz}$), 7.60 (2H, d, $J=9.0\text{Hz}$), 7.89 (1H, d, $J=8.1\text{Hz}$), 8.02 (1H, s)

25 APCI-MASS : $m/z = 417$ ($M^+ + H$)

The following compounds (Preparations 171 to 175) were obtained according to a similar manner to that of Preparation 41.

30

Preparation 171

Ethyl [4-(4-octylphenyl)-2,3-dihydro-4H-1,2,4-triazole-3-one-2-yl]acetate

35 IR (KBr) : 2921.6, 1764.5, 1715, 1197.6 cm^{-1}

NMR (CDCl₃, δ) : 0.88 (3H, t, J=6.7Hz), 1.30 (3H, t, J=7.1Hz), 1.2-1.4 (10H, m), 1.5-1.7 (2H, m), 2.63 (2H, t, J=7.9Hz), 4.26 (2H, q, J=7.1Hz), 4.64 (2H, s), 7.28 (2H, d, J=8.4Hz), 7.44 (2H, d, J=8.4Hz), 7.71 (1H, s)

Preparation 172

4-[4-(4-tert-Butoxycarbonylpiperazin-1-yl)phenyl]-2-(4-methylpentyl)-2,3-dihydro-4H-1,2,4-triazol-3-one

IR (KBr) : 1687.4 cm⁻¹

NMR (CDCl₃, δ) : 0.90 (6H, d, J=6.5Hz), 1.1-1.4 (2H, m), 1.49 (9H, s), 1.4-1.9 (3H, m), 3.16 (4H, t, J=4.9Hz), 3.59 (4H, t, J=4.9Hz), 3.82 (2H, t, J=7.3Hz), 6.98 (2H, d, J=9.0Hz), 7.41 (2H, d, J=9.0Hz), 7.61 (1H, s)

Preparation 173

Methyl 6-(8-bromooctyloxy)-2-naphthoate

IR (KBr) : 2933.2, 2856.1, 1720.2, 1294, 1209.1 cm⁻¹

NMR (CDCl₃, δ) : 1.3-1.6 (8H, m), 1.75-2.0 (4H, m), 3.42 (2H, t, J=6.8Hz), 3.96 (3H, s), 4.09 (2H, t, J=6.5Hz), 7.14 (1H, d, J=1.7Hz), 7.19 (1H, dd, J=8.9 and 1.7Hz), 7.73 (1H, d, J=8.7Hz), 7.83 (1H, d, J=8.9Hz), 8.01 (1H, dd, J=8.7 and 1.7Hz), 8.51 (1H, d, J=1.7Hz)

APCI-MASS : m/z = 393 (M⁺+H)

Preparation 174

4-[4-(6-n-Propyloxyhexyloxy)phenyl]benzoic acid

IR (KBr) : 2937, 2858, 1695, 1683, 1604, 1430, 1290, 1247, 1195 cm⁻¹

NMR (DMSO-d₆, δ) : 0.85 (3H, t, J=7.4Hz), 1.3-1.9 (10H, m), 3.2-3.4 (4H, m), 4.01 (2H, t, J=6.3Hz), 7.04 (2H, d, J=8.7Hz), 7.67 (2H, d, J=8.7Hz), 7.74 (2H, d, J=8.3Hz), 7.98 (2H, d, J=8.3Hz), 12.9 (1H,

s)

APCI-MASS : $m/z = 357$ ($M+H^+$)

Preparation 175

5 4-[4-(6-Bromohexyloxy)phenyl]bromobenzene

NMR ($CDCl_3$, δ) : 1.40-1.65 (4H, m), 1.70-2.00 (4H, m),
3.43 (2H, t, $J=6.7$ Hz), 4.00 (2H, t, $J=6.4$ Hz), 6.95
(2H, d, $J=8.8$ Hz), 7.30-7.60 (6H, m)

10 The following compounds (Preparations 176 to 180) were
obtained according to a similar manner to that of Preparation
43.

Preparation 176

15 4-[4-(4-n-Pentyloxyphenyl)piperazin-1-yl]benzoic acid
dihydrochloride

IR (KBr) : 1668.1, 1602.6, 1510.0, 1228.4 cm^{-1}

20 NMR ($DMSO-d_6$, δ) : 0.89 (3H, t, $J=6.9$ Hz), 1.2-1.5 (5H,
m), 1.6-1.9 (2H, m), 3.0-3.2 (4H, m), 3.4-3.6 (4H,
m), 3.88 (2H, t, $J=6.4$ Hz), 6.83 (2H, d, $J=9$ Hz),
6.9-7.1 (4H, m), 7.79 (2H, d, $J=8.8$ Hz), 12.32 (1H,
s)

APCI-MASS : $m/z = 369$ ($M+H^+$)

25 Preparation 177

4-[4-(4-n-Heptyloxyphenyl)piperazin-1-yl]benzoic acid
dihydrochloride

IR (KBr) : 1666.2, 1600.6, 1511.9 cm^{-1}

30 NMR ($CDCl_3$, δ) : 0.89 (3H, t, $J=6.9$ Hz), 1.2-2.0 (10H,
m), 3.1-3.3 (4H, m), 3.4-3.6 (4H, m), 3.92 (2H, t,
 $J=6.4$ Hz), 6.8-7.1 (6H, m), 8.00 (2H, d, $J=8.8$ Hz)

Preparation 178

35 4-[4-[4-(4-Methylpentyloxy)phenyl]piperazin-1-yl]benzoic

acid dihydrochloride

IR (KBr) : 1668.1, 1602.6, 1510.0, 1236.1 cm^{-1}

NMR (DMSO- d_6 , δ) : 0.89 (6H, d, $J=6.5\text{Hz}$), 1.2-1.4 (2H, m), 1.4-1.8 (3H, m), 3.0-3.2 (4H, m), 3.3-3.5 (4H, m), 3.87 (2H, t, $J=6.3\text{Hz}$), 6.83 (2H, d, $J=9.0\text{Hz}$), 6.9-7.1 (4H, m), 7.79 (2H, d, $J=8.8\text{Hz}$), 12.33 (1H, s)

APCI-MASS : $m/z = 383$ ($M+H^+$)

10 Preparation 179

4-[4-[4-(8-Bromooctyloxy)phenyl]piperazin-1-yl]benzoic acid dihydrochloride

IR (KBr) : 1670.1, 1602.6, 1511.9, 1234.2 cm^{-1}

NMR (DMSO- d_6 , δ) : 1.2-1.5 (8H, m), 1.6-1.9 (4H, m), 3.0-3.2 (4H, m), 3.2-3.5 (4H, m), 3.52 (2H, t, $J=6.7\text{Hz}$), 3.88 (2H, t, $J=6.4\text{Hz}$), 6.83 (2H, d, $J=9.1\text{Hz}$), 6.94 (2H, d, $J=9.1\text{Hz}$), 7.02 (2H, d, $J=8.9\text{Hz}$), 7.79 (2H, d, $J=8.9\text{Hz}$)

20 Preparation 180

3-Fluoro-4-[4-(4-n-hexyloxyphenyl)piperazin-1-yl]benzoic acid dihydrochloride

IR (KBr) : 1673.9, 1511.9, 1240.0 cm^{-1}

NMR (DMSO- d_6 , δ) : 0.88 (3H, t, $J=6.5\text{Hz}$), 1.2-1.5 (6H, m), 1.6-1.8 (2H, m), 3.0-3.5 (8H, m), 3.88 (2H, t, $J=6.4\text{Hz}$), 6.7-7.2 (5H, m), 7.4-7.8 (2H, m), 12.82 (1H, s)

APCI-MASS : $m/z = 401$ (M^++H)

30 The following compound was obtained according to a similar manner to that of Preparation 46.

Preparation 181

35 1-(4-Methoxycarbonylphenyl)-3-(4-n-hexyloxyphenyl)-propan-1,3-dione

IR (KBr) : 2956, 2927, 2856, 1722, 1511, 1284,
1108 cm^{-1}

NMR (CDCl_3 , δ) : 0.92 (3H, t, $J=6.4\text{Hz}$), 1.2-2.0 (8H,
m), 3.96 (3H, s), 4.04 (2H, t, $J=6.5\text{Hz}$), 6.82 (1H,
s), 6.97 (2H, d, $J=8.7\text{Hz}$), 7.9-8.1 (4H, m), 8.14
(2H, d, $J=8.3\text{Hz}$)

APCI-MASS : $m/z = 383$ ($\text{M}+\text{H}^+$)

The following compounds (Preparations 182 to 185) were
obtained according to a similar manner to that of Preparation
47.

Preparation 182

Methyl 5-(4-octyloxyphenyl)-1-methylpyrazole-3-
carboxylate

IR (KBr pelet) : 2923, 1724, 1616, 1513, 1446, 1251,
1120 cm^{-1}

NMR (CDCl_3 , δ) : 0.89 (3H, t, $J=6.8\text{Hz}$), 1.2-1.5 (10H,
m), 1.7-1.9 (2H, m), 3.90 (3H, s), 3.98 (2H, t,
 $J=6.6\text{Hz}$), 4.20 (3H, s), 6.92 (2H, d, $J=8.9\text{Hz}$), 7.04
(1H, s), 7.89 (2H, d, $J=8.9\text{Hz}$)

APCI-MASS : $m/z = 345$ ($\text{M}+\text{H}^+$)

Preparation 183

Methyl 4-[5-(4-n-pentyloxyphenyl)pyrazol-3-yl]benzoate

IR (KBr) : 3236, 2952, 2873, 1716, 1616, 1508, 1276,
1174, 1106 cm^{-1}

NMR (CDCl_3 , δ) : 0.94 (3H, t, $J=7.0\text{Hz}$), 1.3-1.5 (4H,
m), 1.7-1.9 (2H, m), 3.92 (3H, s), 3.96 (2H, t,
 $J=6.7\text{Hz}$), 6.78 (1H, s), 6.88 (2H, d, $J=8.7\text{Hz}$), 7.55
(2H, d, $J=8.7\text{Hz}$), 7.79 (2H, d, $J=8.4\text{Hz}$), 8.02 (2H,
d, $J=8.4\text{Hz}$)

APCI-MASS : $m/z = 365$ ($\text{M}+\text{H}^+$)

Preparation 184

Methyl 5-(4-octyloxyphenyl)isoxazole-3-carboxylate

IR (KBr pelet) : 2950, 2921, 1724, 1614, 1510, 1446,
1257, 1178, 1143, 1009 cm^{-1}

5 NMR (CDCl_3 , δ) : 0.89 (3H, t, $J=6.8\text{Hz}$), 1.2-1.6 (10H, m), 1.7-1.9 (2H, m), 4.0-4.1 (5H, m), 6.80 (1H, s), 6.98 (2H, dd, $J=6.9$ and 2.1Hz), 7.73 (2H, dd, $J=6.9$ and 2.1Hz)

APCI-MASS : $m/z = 332$ ($M+H^+$)

10 Preparation 185

Methyl 4-[3-(4-n-hexyloxyphenyl)pyrazol-5-yl]benzoate

IR (KBr) : 2952, 1716, 1616, 1508, 1276, 1106 cm^{-1}

15 NMR (CDCl_3 , δ) : 0.91 (3H, t, $J=6.3\text{Hz}$), 1.2-1.6 (6H, m), 1.7-1.9 (2H, m), 3.8-4.0 (5H, m), 6.76 (1H, s), 6.86 (2H, d, $J=8.8\text{Hz}$), 7.54 (2H, d, $J=8.8\text{Hz}$), 7.77 (2H, d, $J=8.4\text{Hz}$), 8.00 (2H, d, $J=8.4\text{Hz}$)

APCI-MASS : $m/z = 379$ ($M+H^+$)

Preparation 186

20 A suspension of 1-(4-n-Pentyloxyphenyl)-3-(4-ethoxycarbonylphenyl)-1-buten-3-one (74.43 g) and hydroxyamine hydrochloride (28.23 g) and potassium carbonate (56.11 g) in ethanol (400 ml) was refluxed for 4 hours. The mixture was diluted with ethyl acetate, washed with water
25 (x 2), brine and dried over magnesium sulfate. The solvents were removed under reduced pressure to give crude oxime. To a solution of crude oxime in dichloroethane (500 ml) was added activated-manganese(IV) oxide (200 g). The reaction mixture was refluxed for 2 hours and filtered. The residue
30 was washed with dichloromethane. The solvents were removed under reduced pressure and the residue was triturated with acetonitrile. The solid was collected by filtration and dried to give ethyl 4-[5-(4-n-Pentyloxyphenyl)isoxazol-3-yl]benzoate (21.07 g).

35 IR (KBr) : 2945, 2872, 1717, 1615, 1508, 1280,

1108 cm^{-1}

NMR (CDCl_3 , δ) : 0.95 (3H, t, $J=6.9\text{Hz}$), 1.3-1.9 (9H, m), 4.01 (2H, t, $J=6.5\text{Hz}$), 4.41 (2H, q, $J=7.1\text{Hz}$), 6.74 (1H, s), 6.99 (2H, d, $J=8.8\text{Hz}$), 7.76 (2H, d, $J=8.8\text{Hz}$), 7.93 (2H, d, $J=8.4\text{Hz}$), 8.15 (2H, d, $J=8.4\text{Hz}$)

APCI-MASS : $m/z = 380$ ($\text{M}+\text{H}^+$)

The following compounds (Preparations 187 to 190) were obtained according to a similar manner to that of Preparation 48.

Preparation 187

Methyl 6-[4-[4-(8-Methoxyoctyloxy)phenyl]piperazin-1-yl]nicotinate

IR (KBr) : 2933, 2858, 1722, 1608, 1513, 1432, 1405, 1278, 1245 cm^{-1}

NMR (CDCl_3 , δ) : 1.3-1.9 (12H, m), 3.16 (4H, t, $J=5.0\text{Hz}$), 3.33 (3H, s), 3.36 (2H, t, $J=6.5\text{Hz}$), 3.8-4.0 (9H, m), 6.64 (1H, d, $J=9.1\text{Hz}$), 6.85 (2H, d, $J=9.2\text{Hz}$), 6.93 (2H, d, $J=9.2\text{Hz}$), 8.04 (1H, dd, $J=9.1$ and 2.2Hz), 8.81 (1H, d, $J=2.2\text{Hz}$)

APCI-MASS : $m/z = 456$ ($\text{M}+\text{H}^+$)

Preparation 188

4-[4-(5-Methoxypentyloxy)phenyl]bromobenzene

IR (KBr) : 2940, 2856, 1604, 1479, 1286, 1255, 1124 cm^{-1}

NMR (CDCl_3 , δ) : 1.5-1.9 (6H, m), 3.34 (3H, s), 3.41 (2H, t, $J=6.1\text{Hz}$), 3.99 (2H, t, $J=6.4\text{Hz}$), 6.95 (2H, d, $J=8.7\text{Hz}$), 7.4-7.6 (6H, m)

APCI-MASS : $m/z = 349$ ($\text{M}+\text{H}^+$)

Preparation 189

Methyl 6-(8-methoxyoctyloxy)-2-naphthoate

NMR (DMSO-d₆, δ) : 1.2-1.6 (10H, m), 1.7-1.9 (2H, m),
3.20 (3H, s), 3.29 (2H, t, J=6.4Hz), 3.89 (3H, s),
4.11 (2H, t, J=6.4Hz), 7.24 (1H, dd, J=9.0 and
2.4Hz), 7.40 (1H, d, J=2.4Hz), 7.88 (1H, d,
5 J=8.7Hz), 7.94 (1H, dd, J=8.7 and 1.5Hz), 8.03 (1H,
d, J=9.0Hz), 8.55 (1H, d, J=1.5Hz)

Preparation 190

4-[4-[4-(8-Methoxyoctyloxy)phenyl]piperazin-1-yl]benzoic
10 acid dihydrochloride

IR (KBr) : 1668.1, 1602.6, 1511.9, 1236.1 cm⁻¹

NMR (DMSO-d₆, δ) : 1.2-1.8 (12H, m), 3.05-3.2 (4H, m),
3.29 (2H, t, J=7.1Hz), 3.33 (3H, s), 3.4-3.55 (4H,
m), 3.88 (2H, t, J=6.4Hz), 6.82 (2H, d, J=9.0Hz),
15 6.94 (2H, d, J=9.0Hz), 7.02 (2H, d, J=8.8Hz), 7.79
(2H, d, J=8.8Hz), 12.31 (1H, s)

The following compounds (Preparations 191 to 254) were
obtained according to a similar manner to that of Preparation
20 49.

Preparation 191

1-[4-[4-[4-[2-(4-Methylpentyl)-2,3-dihydro-4H-1,2,4-
triazol-3-one-4-yl]phenyl]piperazin-1-yl]benzoyl]-
25 benzotriazole 3-oxide

IR (KBr) : 1766.5, 1693.2, 1600.6, 1519.6 cm⁻¹

Preparation 192

1-[4-(4-Octylphenyl)-2,3-dihydro-4H-1,2,4-triazol-3-one-
30 2-yl-acetyl]benzotriazole 3-oxide

IR (KBr) : 2921.6, 1753.0, 1720.0, 1423.2 cm⁻¹

NMR (CDCl₃, δ) : 0.88 (3H, t, J=6.7Hz), 1.2-1.4 (10H,
m), 1.5-1.8 (2H, m), 2.65 (2H, t, J=7.5Hz), 5.46
(2H, s), 7.30 (2H, d, J=8.5Hz), 7.48 (2H, d,
35 J=8.5Hz), 7.62 (1H, t, J=8.3Hz), 7.80 (1H, s), 7.82

(1H, t, J=8.3Hz), 8.05 (1H, d, J=8.3Hz), 8.37 (1H, d, J=8.3Hz)

Preparation 193

1-[4-[4-[4-(7-Methoxyheptyloxy)phenyl]piperazin-1-yl]benzoyl]benzotriazole 3-oxide

IR (KBr) : 1783.8, 1600.6, 1511.9, 1232.3, 1184.1 cm^{-1}

NMR (CDCl_3 , δ) : 1.3-1.9 (10H, m), 3.2-3.3 (4H, m), 3.34 (3H, s), 3.38 (2H, t, J=6.4Hz), 3.5-3.7 (4H, m), 3.92 (2H, t, J=6.5Hz), 6.87 (2H, d, J=9.2Hz), 6.95 (2H, d, J=9.2Hz), 7.00 (2H, d, J=9.0Hz), 7.3-7.6 (3H, m), 8.09 (1H, d, J=8.2Hz), 8.15 (2H, d, J=9.0Hz)

Preparation 194

1-[4-[4-(4-n-Heptyloxyphenyl)piperazin-1-yl]benzoyl]benzotriazole 3-oxide

IR (KBr) : 1783.8, 1600.6, 1511.9, 1230.4, 1184.1 cm^{-1}

NMR (CDCl_3 , δ) : 0.90 (3H, t, J=6.3Hz), 1.2-1.6 (8H, m), 1.7-1.9 (2H, m), 3.2-3.3 (4H, m), 3.5-3.7 (4H, m), 3.93 (2H, t, J=6.5Hz), 6.87 (2H, d, J=9.2Hz), 6.95 (2H, d, J=9.2Hz), 7.00 (2H, d, J=9.0Hz), 7.3-7.7 (3H, m), 8.09 (1H, d, J=8.2Hz), 8.15 (2H, d, J=9.0Hz)

Preparation 195

1-[4-[4-[4-(4-Methylpentyloxy)phenyl]piperazin-1-yl]benzoyl]benzotriazole 3-oxide

NMR (CDCl_3 , δ) : 0.92 (6H, d, J=6.6Hz), 1.2-1.4 (2H, m), 1.5-1.9 (3H, m), 3.1-3.3 (4H, m), 3.5-3.7 (4H, m), 3.92 (2H, t, J=6.6Hz), 6.87 (2H, d, J=9.3Hz), 6.96 (2H, d, J=9.3Hz), 7.01 (2H, d, J=9.0Hz), 7.4-7.6 (3H, m), 8.10 (1H, d, J=8.2Hz), 8.15 (2H, d, J=9.0Hz)

Preparation 196

1-[4-[4-(4-n-Pentyloxyphenyl)piperazin-1-yl]benzoyl]benzotriazole 3-oxide

IR (KBr) : 1787.7, 1600.6, 1511.9, 1232.3, 1184.1 cm^{-1}

5 NMR (CDCl_3 , δ) : 0.93 (3H, t, $J=6.9\text{Hz}$), 1.3-1.6 (4H, m), 1.7-1.9 (2H, m), 3.1-3.4 (4H, m), 3.5-3.8 (4H, m), 3.93 (2H, t, $J=6.6\text{Hz}$), 6.87 (2H, d, $J=9.2\text{Hz}$), 6.92 (2H, d, $J=9.2\text{Hz}$), 7.01 (2H, d, $J=9.1\text{Hz}$), 7.4-7.6 (3H, m), 8.10 (1H, d, $J=8.2\text{Hz}$), 8.15 (2H, d, $J=9.1\text{Hz}$)

Preparation 197

1-[4-[4-[8-(1H-Tetrazol-1-yl)octyloxy]phenyl]benzoyl]-benzotriazole 3-oxide

15 and

1-[4-[4-[8-(2H-tetrazol-2-yl)octyloxy]phenyl]benzoyl]-benzotriazole 3-oxide

20 IR (KBr) : 1778.0, 1602.6, 1189.9, 981.6 cm^{-1}

NMR (CDCl_3 , δ) : 1.2-1.6 (8H, m), 1.7-1.9 (2H, m), 1.9-2.2 (2H, m), 4.02 (2H, t, $J=6.4\text{Hz}$), 4.44 and 4.66 (2H, t, $J=7.1\text{Hz}$), 7.02 (2H, d, $J=8.8\text{Hz}$), 7.4-7.6 (3H, m), 7.63 (2H, d, $J=8.8\text{Hz}$), 7.79 (2H, d, $J=8.6\text{Hz}$), 8.12 (1H, d, $J=8.2\text{Hz}$), 8.32 (2H, d, $J=8.6\text{Hz}$), 8.51 and 8.60 (1H, s)

Preparation 198

30 1-[4-[4-[8-(2,6-Dimethylmorpholin-4-yl)octyloxy]phenyl]benzoyl]benzotriazole 3-oxide

IR (KBr) : 1778.0, 1600.6, 977.7 cm^{-1}

35 NMR (CDCl_3 , δ) : 1.18 (6H, d, $J=6.3\text{Hz}$), 1.2-1.7 (10H, m), 1.7-2.0 (4H, m), 2.4-2.6 (2H, m), 2.9-3.2 (2H, m), 3.7-3.9 (2H, m), 4.01 (2H, t, $J=6.5\text{Hz}$), 7.02 (2H, d, $J=8.8\text{Hz}$), 7.4-7.7 (3H, m), 7.63 (2H, d,

J=8.8Hz), 7.79 (2H, d, J=8.5Hz), 8.12 (1H, d, J=8.1Hz), 8.32 (2H, d, J=8.5Hz)

Preparation 199

5 1-[6-[4-(4-Octyloxyphenyl)piperazin-1-yl]nicotinoyl]-
benzotriazole 3-oxide

IR (KBr pelet) : 2922, 2854, 1766, 1602, 1513, 1417,
1234, 1025, 950, 813 cm^{-1}

10 NMR (CDCl_3 , δ) : 0.89 (3H, t, J=6.8Hz), 1.2-1.5 (10H,
m), 1.7-1.9 (2H, m), 3.1-3.3 (4H, m), 3.9-4.1 (6H,
m), 6.75 (1H, d, J=9.2Hz), 6.87 (2H, d, J=9.2Hz),
6.95 (2H, d, J=9.2Hz), 7.4-7.6 (3H, m), 8.10 (1H,
d, J=8.1Hz), 8.19 (1H, dd, J=9.2 and 2.4Hz), 9.04
(1H, d, J=2.4Hz)

15 APCI-MASS : m/z = 529 ($\text{M}+\text{H}^+$)

Preparation 200

1-[2-(4-Hexyloxyphenyl)benzoxazol-5-yl-carbonyl]-
benzotriazole 3-oxide

20 IR (KBr) : 2950, 1774, 1623, 1504, 1265, 1176 cm^{-1}

NMR (CDCl_3 , δ) : 0.93 (3H, t, J=6.9Hz), 1.3-1.6 (6H,
m), 1.8-2.0 (2H, m), 4.07 (2H, t, J=6.5Hz), 7.06
(2H, d, J=8.9Hz), 7.4-7.6 (3H, m), 7.75 (1H, d,
J=8.6Hz), 8.13 (1H, d, J=8.2Hz), 8.2-8.4 (3H, m),
25 8.67 (1H, d, J=1.6Hz)

APCI-MASS : m/z = 457 ($\text{M}+\text{H}^+$)

Preparation 201

30 1-[4-[4-(4-n-Butyloxyphenyl)phenyl]benzoyl]-
benzotriazole 3-oxide

IR (KBr) : 2958, 2871, 1776, 1600, 1398, 1255, 1211,
1037 cm^{-1}

35 NMR (CDCl_3 , δ) : 1.00 (3H, t, J=7.2Hz), 1.4-1.9 (4H,
m), 4.03 (2H, t, J=6.4Hz), 7.01 (2H, d, J=8.3Hz),
7.4-7.8 (9H, m), 7.87 (2H, d, J=8.1Hz), 8.12 (1H,

d, J=8.4Hz), 8.36 (2H, d, J=7.9Hz)
APCI-MASS : m/z = 464 (M+H)⁺

Preparation 202

5 1-[2-(4-Heptyloxyphenyl)pyridin-5-yl-
carbonyl]benzotriazole 3-oxide

IR (KBr) : 2944, 2867, 1793, 1770, 1589, 1471, 1321,
1093 cm⁻¹

10 NMR (CDCl₃, δ) : 0.91 (3H, t, J=6.7Hz), 1.2-1.6 (8H,
m), 1.7-1.9 (2H, m), 4.05 (2H, t, J=6.5Hz), 7.04
(2H, d, J=8.0Hz), 7.4-7.6 (3H, m), 7.91 (1H, d,
J=8.5Hz), 8.1-8.2 (3H, m), 8.51 (1H, dd, J=8.5 and
2.3Hz), 9.47 (1H, d, J=2.3Hz)

APCI-MASS : m/z = 431 (M+H)⁺

15 Preparation 203

1-[2-(2-Octyloxyppyridin-5-yl)benzoxazol-5-yl-
carbonyl]benzotriazole 3-oxide

IR (KBr pelet) : 2925, 2854, 1787, 1623, 1479, 1263,
989 cm⁻¹

20 NMR (CDCl₃, δ) : 0.89 (3H, t, J=6.8Hz), 1.2-1.5 (10H,
m), 1.8-1.9 (2H, m), 4.42 (2H, t, J=6.7Hz), 6.91
(1H, d, J=8.7Hz), 6.4-6.6 (3H, m), 7.79 (1H, d,
J=8.6Hz), 8.13 (1H, d, J=8.2Hz), 8.32 (1H, dd,
25 J=8.6 and 1.7Hz), 8.41 (1H, dd, J=8.7 and 2.4Hz),
8.70 (1H, d, J=1.4Hz), 9.07 (1H, d, J=1.9Hz)

APCI-MASS : m/z = 486 (M+H)⁺

Preparation 204

30 1-[2-[4-(4-Hexylphenyl)phenyl]benzoxazol-5-yl-
carbonyl]benzotriazole 3-oxide

IR (KBr) : 2927, 2854, 1785, 1621, 1490, 1261, 1166,
1052 cm⁻¹

35 NMR (CDCl₃, δ) : 0.90 (3H, t, J=6.5Hz), 1.2-1.8 (8H,
m), 2.68 (2H, t, J=7.9Hz), 7.31 (2H, d, J=8.2Hz),

7.4-7.7 (5H, m), 7.79-7.81 (3H, m), 8.13 (1H, d, J=8.3Hz), 8.3-8.4 (3H, m), 8.73 (1H, d, J=1.3Hz)

APCI-MASS : m/z = 517 (M+H⁺)

5 Preparation 205

1-[2-[4-(4-n-Butyloxyphenyl)phenyl]pyridin-5-yl-carbonyl]benzotriazole 3-oxide

IR (KBr) : 2956, 2933, 2871, 1774, 1650, 1591, 1471, 1251 cm⁻¹

10 NMR (CDCl₃, δ) : 1.00 (3H, t, J=7.2Hz), 1.5-1.9 (4H, m), 4.03 (2H, t, J=6.4Hz), 7.02 (2H, d, J=8.6Hz), 7.4-7.6 (3H, m), 7.54 (2H, d, J=7.3Hz), 7.62 (2H, d, J=8.5Hz), 8.02 (1H, d, J=8.3Hz), 8.13 (1H, d, J=8.2Hz), 8.21 (2H, d, J=7.9Hz), 8.57 (1H, dd, J=8.3 and 2.0Hz), 9.54 (1H, d, J=2.0Hz)

15 APCI-MASS : m/z = 465 (M+H)⁺

Preparation 206

20 1-[4-[4-(5-Phenoxypropyloxy)phenyl]benzoyl]-benzotriazole 3-oxide

IR (KBr) : 2944, 2869, 1770, 1600, 1494, 1249, 1189 cm⁻¹

25 NMR (CDCl₃, δ) : 1.6-1.8 (2H, m), 1.8-2.0 (4H, m), 4.01 (2H, t, J=6.3Hz), 4.07 (2H, t, J=6.2Hz), 6.91 (2H, d, J=8.9Hz), 7.04 (2H, d, J=8.7Hz), 7.3-7.6 (4H, m), 7.63 (2H, d, J=8.6Hz), 7.78 (2H, d, J=8.4Hz), 8.12 (1H, d, J=8.1Hz), 8.32 (2H, d, J=8.4Hz)

APCI-MASS : m/z = 494 (M+H)⁺

30

Preparation 207

1-[4-[5-(4-Hexyloxyphenyl)-1,3,4-oxadiazol-2-yl]benzoyl]benzotriazole 3-oxide

35 IR (KBr) : 2956, 2921, 2856, 1778, 1612, 1496, 1261, 1232, 1025 cm⁻¹

NMR (CDCl₃, δ) : 0.92 (3H, t, J=6.7Hz), 1.3-1.6 (6H, m), 1.8-2.0 (2H, m), 4.05 (2H, t, J=6.5Hz), 7.05 (2H, d, J=8.7Hz), 7.4-7.6 (3H, m), 8.10 (2H, d, J=8.7Hz), 8.13 (1H, d, J=7.4Hz), 8.37 (2H, d, J=8.5Hz), 8.45 (2H, d, J=8.5Hz)

APCI-MASS : m/z = 484 (M+H)⁺

Preparation 208

1-[4-[5-(4-n-Hexyloxyphenyl)-1,3,4-thiadiazol-2-yl]benzoyl]benzotriazole 3-oxide

IR (KBr) : 2952, 2873, 1774, 1602, 1261, 1230, 1176 cm⁻¹

NMR (CDCl₃, δ) : 0.93 (3H, t, J=6.8Hz), 1.3-2.0 (8H, m), 4.04 (2H, t, J=6.5Hz), 7.02 (2H, d, J=8.7Hz), 7.4-7.7 (3H, m), 7.98 (2H, d, J=8.7Hz), 8.13 (1H, d, J=8.7Hz), 8.25 (2H, d, J=8.3Hz), 8.41 (2H, d, J=8.3Hz)

APCI-MASS : m/z = 500 (M+H)⁺

Preparation 209

1-[5-(4-Octyloxyphenyl)-1-methylpyrazol-3-yl-carbonyl]benzotriazole 3-oxide

IR (KBr pelet) : 2939, 2852, 1776, 1687, 1612, 1448, 1249, 995 cm⁻¹

NMR (CDCl₃, δ) : 0.89 (3H, t, J=6.7Hz), 1.3-1.5 (10H, m), 1.7-1.9 (2H, m), 4.01 (2H, t, J=6.5Hz), 4.25 (3H, s), 6.97 (2H, d, J=6.8Hz), 7.4-7.7 (4H, m), 7.78 (2H, d, J=6.8Hz), 8.14 (1H, d, J=8.0Hz)

APCI-MASS : m/z = 448 (M+H)⁺

Preparation 210

1-[4-[5-(4-n-Pentyloxyphenyl)pyrazol-3-yl]benzoyl]benzotriazole 3-oxide

IR (KBr) : 3251, 2956, 2869, 1780, 1612, 1506, 1232, 985 cm⁻¹

NMR (CDCl₃, δ) : 0.95 (3H, t, J=6.9Hz), 1.3-1.6 (4H, m), 1.7-2.0 (2H, m), 4.01 (2H, t, J=6.6Hz), 6.90 (1H, s), 6.99 (2H, d, J=8.7Hz), 7.4-7.6 (5H, m), 8.0-8.2 (3H, m), 8.33 (2H, d, J=8.4Hz)

5 APCI-MASS : m/z = 468 (M+H⁺)

Preparation 211

1-[5-[4-(4-n-Butoxyphenyl)phenyl]furan-2-yl-carbonyl]benzotriazole 3-oxide

10 IR (KBr) : 2958, 2871, 1781, 1678, 1603, 1535, 1479, 1265 cm⁻¹

NMR (CDCl₃, δ) : 1.00 (3H, t, J=7.3Hz), 1.4-1.9 (4H, m), 4.02 (2H, t, J=6.4Hz), 6.9-7.1 (3H, m), 7.4-8.2 (11H, m)

15 APCI-MASS : m/z = 351 (Methyl ester)

Preparation 212

1-(3-(S)-Hydroxy-2-benzylhexadecanoyl)benzotriazole 3-oxide

20 IR (Neat) : 2854.1, 1814.7, 1459.8, 742.5 cm⁻¹

Preparation 213

1-(3-(R)-Benzyloxycarboxylamino-18-methoxyoctadecanoyl)-benzotriazole 3-oxide

25 IR (KBr) : 1805.0, 1729.8, 1695.1 cm⁻¹

NMR (DMSO-d₆, δ) : 1.1-1.65 (30H, m), 3.20 (3H, s), 3.28 (2H, t, J=6.5Hz), 4.01 (1H, m), 5.06 (2H, s), 7.32 (5H, m), 7.4-7.8 (3H, m), 8.12 (1H, d, J=7Hz)

30 Preparation 214

1-(3-(S)-Hydroxyhexadecanoyl)benzotriazole 3-oxide

IR (KBr) : 1710.6, 1498.4, 1429.0, 771.4 cm⁻¹

35 NMR (CDCl₃, δ) : 0.88 (3H, t, J=6.4Hz), 1.2-1.7 (24H, m), 2.00 (1H, s), 3.1-3.5 (2H, m), 4.30 (1H, m), 7.59 (1H, t, J=7.8Hz), 7.81 (1H, t, J=7.8Hz), 8.02

(1H, d, J=8.3Hz), 8.42 1(1H, d, J=8.3Hz)

Preparation 215

1-(3-Methyl-2-tridecenoyl)benzotriazole 3-oxide

5 IR (KBr) : 2927.4, 1791.5, 1633.4, 1081.9 cm^{-1}

NMR (CDCl_3 , δ) : 0.89 (3H, t, J=6.3Hz), 1.1-1.7 (20H, m), 2.25 (3H, s), 6.08 (1H, s), 7.3-7.6 (3H, m), 8.06 (1H, d, J=8.2Hz)

10 Preparation 216

1-[4-[4-[4-(8-Methoxyoctyloxy)phenyl]piperazin-1-yl]benzoyl]benzotriazole 3-oxide

IR (KBr) : 1780.0, 1600.6, 1511.9, 1234.2, 1184.1 cm^{-1}

15 NMR (CDCl_3 , δ) : 1.3-1.9 (12H, m), 3.24 (4H, t, J=5.0Hz), 3.33 (3H, s), 3.37 (2H, t, J=6.8Hz), 3.62 (4H, t, J=5.0Hz), 3.92 (2H, t, J=6.5Hz), 6.8-7.1 (6H, m), 7.35-7.65 (3H, m), 8.09 (1H, d, J=8.2Hz), 8.15 (2H, d, J=9.0Hz)

20 Preparation 217

1-[3-Fluoro-4-[4-(4-n-hexyloxyphenyl)piperazin-1-yl]benzoyl]benzotriazole 3-oxide

IR (KBr) : 1778.0 cm^{-1}

25 Preparation 218

1-[3-Chloro-4-[4-(4-n-hexyloxyphenyl)piperazin-1-yl]benzoyl]benzotriazole 3-oxide

IR (KBr) : 1778.0, 1594.8, 1511.9, 1218.8 cm^{-1}

30 NMR (CDCl_3 , δ) : 0.91 (3H, t, J=6.5Hz), 1.2-1.6 (6H, m), 1.6-1.9 (2H, m), 3.29 (4H, t, J=3.6Hz), 3.44 (4H, t, J=3.6Hz), 3.93 (2H, t, J=6.5Hz), 6.87 (2H, d, J=9.2Hz), 6.97 (2H, d, J=9.2Hz), 7.19 (1H, d, J=8.6Hz), 7.4-7.7 (3H, m), 8.10 (1H, d, J=6.4Hz), 8.14 (1H, dd, J=8.6 and 2.1Hz), 8.27 (1H, d, J=2.1Hz)

35

467250" E2260880

APCI-MASS : $m/z = 534$ ($M^+ + H$)

Preparation 219

1-[4-(4-Piperidinopiperidin-1-yl)benzoyl]benzotriazole
3-oxide

IR (KBr) : 1758.8, 1602.6, 1186.0 cm^{-1}

NMR (CDCl_3 , δ) : 1.35-1.8 (8H, m), 1.96 (2H, d, $J=13\text{Hz}$), 2.45-2.7 (5H, m), 2.97 (2H, td, $J=12.8$ and 2.6Hz), 4.04 (2H, d, $J=13\text{Hz}$), 6.93 (2H, d, $J=9.2\text{Hz}$), 7.35-7.6 (3H, m), 8.1-8.4 (3H, m)

Preparation 220

1-[3-[4-(4-n-Hexyloxyphenyl)piperazin-1-yl]pyridazin-6-yl-carbonyl]benzotriazole 3-oxide

IR (KBr) : 1787.7, 1585.2, 1511.9, 1240.0 cm^{-1}

Preparation 221

1-[5-[4-(4-n-Hexyloxyphenyl)piperazin-1-yl]picolinoyl]-benzotriazole 3-oxide

IR (KBr) : 1766.5, 1575.6, 1511.9, 1232.3 cm^{-1}

NMR (CDCl_3 , δ) : 0.91 (3H, t, $J=6.5\text{Hz}$), 1.2-1.6 (6H, m), 1.65-1.9 (2H, m), 3.27 (4H, t, $J=5.1\text{Hz}$), 3.66 (4H, t, $J=5.1\text{Hz}$), 3.93 (2H, t, $J=6.5\text{Hz}$), 6.88 (2H, d, $J=9.2\text{Hz}$), 6.95 (2H, d, $J=9.2\text{Hz}$), 7.25 (1H, dd, $J=7.6$ and 2.9Hz), 7.35-7.6 (3H, m), 8.09 (1H, d, $J=8.2\text{Hz}$), 8.18 (1H, d, $J=8.9\text{Hz}$), 8.52 (1H, d, $J=2.9\text{Hz}$)

APCI-MASS : $m/z = 501$ ($M^+ + H$)

Preparation 222

1-[4-[4-(4-Cyclohexylphenyl)piperazin-1-yl]benzoyl]-benzotriazole 3-oxide

IR (KBr) : 1770.3, 1602.6, 1515.8, 1232.3, 1186.0 cm^{-1}

NMR (CDCl_3 , δ) : 1.15-1.5 (6H, m), 1.65-2.0 (4H, m), 2.45 (1H, m), 3.33 (4H, t, $J=5.1\text{Hz}$), 3.62 (4H, t,

J=5.1Hz), 6.92 (2H, d, J=8.7Hz), 6.99 (2H, d, J=9.2Hz), 7.16 (2H, d, J=8.7Hz), 7.35-7.65 (3H, m), 8.09 (1H, d, J=8.2Hz), 8.15 (2H, d, J=9.2Hz)

5 Preparation 223

1-[4-[4-(4-n-Hexylphenyl)piperazin-1-yl]benzoyl]-
benzotriazole 3-oxide

IR (KBr) : 1768.4, 1602.6, 1515.8, 1230.4, 1184.1 cm^{-1}

10 NMR (CDCl_3 , δ) : 0.89 (3H, t, J=6.5Hz), 1.2-1.45 (6H, m), 1.5-1.7 (2H, m), 2.55 (2H, t, J=7.6Hz), 3.2-3.4 (4H, m), 3.5-3.7 (4H, m), 6.91 (2H, d, J=8.6Hz), 7.00 (2H, d, J=9.1Hz), 7.13 (2H, d, J=8.5Hz), 7.35-7.6 (3H, m), 8.09 (1H, d, J=8.2Hz), 8.15 (2H, d, J=9.1Hz)

15

Preparation 224

1-[4-[4-(4-Phenylcyclohexyl)piperazin-1-yl]benzoyl]benzotriazole 3-oxide

IR (KBr) : 1780.0, 1762.6, 1602.6, 1234.2, 1182.2 cm^{-1}

20 NMR (CDCl_3 , δ) : 1.3-1.7 (4H, m), 1.95-2.15 (4H, m), 2.35-2.6 (2H, m), 2.79 (4H, t, J=5.0Hz), 3.49 (4H, t, J=5.0Hz), 6.95 (2H, d, J=9.0Hz), 7.1-7.35 (5H, m), 7.35-7.6 (3H, m), 8.08 (1H, d, J=7.1Hz), 8.12 (2H, d, J=9.0Hz)

25

Preparation 225

1-[4-[4-[1-(4-n-Hexyloxyphenyl)piperidin-4-yl]piperazin-1-yl]benzoyl]benzotriazole 3-oxide

IR (KBr) : 1768.4, 1602.6, 1511.9, 1234.2 cm^{-1}

30 NMR (CDCl_3 , δ) : 0.90 (3H, t, J=6.5Hz), 1.2-1.55 (6H, m), 1.6-1.9 (4H, m), 1.96 (2H, d, J=11Hz), 2.44 (1H, m), 2.64 (2H, d, J=1.1Hz), 2.77 (4H, t, J=5.0Hz), 3.48 (4H, t, J=5.0Hz), 3.59 (2H, d, J=11Hz), 3.91 (2H, t, J=6.5Hz), 6.7-7.05 (6H, m), 35 7.35-7.6 (3H, m), 8.08 (1H, d, J=6.9Hz), 8.12 (2H,

d, $J=7.7\text{Hz}$)

Preparation 226

1-[4-(4-Trans-n-pentylcyclohexyl)benzoyl]benzotriazole
5 3-oxide

IR (KBr) : 1799.3, 1778.0, 1608.3, 1228.4, 977.7 cm^{-1}

NMR (CDCl_3 , δ) : 0.91 (3H, t, $J=6.6\text{Hz}$), 1.0-1.7 (13H,
m), 1.93 (4H, d, $J=9.8\text{Hz}$), 2.62 (1H, t, $J=12\text{Hz}$),
7.35-7.6 (5H, m), 8.09 (1H, d, $J=7.9\text{Hz}$), 8.19 (2H,
10 d, $J=8.4\text{Hz}$)

Preparation 227

1-[6-(8-Methoxyoctyloxy)-2-naphthoyl]benzotriazole 3-
oxide

15 IR (KBr) : 2931.3, 2856.1, 1778.0, 1623.8 cm^{-1}

Preparation 228

1-(E)-[3-[4-[4-(7-Fluoroheptyloxy)phenyl]phenyl]-
20 acryloyl]benzotriazole 3-oxide

IR (KBr) : 3070.1, 2935.1, 2859.9, 1700.9, 1619.9,
1596.8 cm^{-1}

NMR (CDCl_3 , δ) : 1.30-2.00 (10H, m), 4.02 (2H, t,
 $J=6.4\text{Hz}$), 4.45 (2H, dt, $J=47.5$ and 6.2Hz), 6.70-
8.65 (14H, m)

25

Preparation 229

1-(6-Heptylnaphthalene-2-carbonyl)benzotriazole 3-oxide

NMR ($\text{DMSO}-d_6$, δ) : 0.75-0.93 (3H, m), 1.10-1.45 (8H,
m), 1.55-1.80 (2H, m), 2.68-2.90 (2H, m), 7.35-9.06
30 (10H, m)

APCI-MASS : $m/z = 388$ (M^++1)

Preparation 230

1-(E)-[3-[4-[4-(8-Methoxyoctyloxy)phenyl]phenyl]-
35 acryloyl]benzotriazole 3-oxide

Preparation 231

1-(E)-[3-[4-[4-(5-Hexenyloxy)phenyl]phenyl]acryloyl]-
benzotriazole 3-oxide

IR (KBr) : 3072.0, 3033.5, 2939.0, 2865.7, 1780.0,
1693.2, 1619.9, 1596.8 cm^{-1}

NMR (DMSO- d_6 , δ) : 1.43-1.66 (2H, m), 1.66-1.90 (2H,
m), 2.02-2.23 (2H, m), 3.90-4.16 (2H, m), 4.90-5.13
(2H, m), 5.72-6.00 (1H, m), 6.93-8.30 (14H, m)

APCI-MASS : m/z = 337 (Methyl ester, $\text{M}^+ + 1$)

Preparation 232

1-(E)-[3-[4-[4-(4-Methylpentyloxy)phenyl]phenyl]-
acryloyl]benzotriazole 3-oxide

IR (KBr) : 3072.0, 3033.5, 2952.5, 2869.6, 1780.0,
1693.2, 1618.0, 1598.7 cm^{-1}

NMR (DMSO- d_6 , δ) : 0.90 (6H, d, $J=6.5\text{Hz}$), 1.20-1.40
(2H, m), 1.50-1.90 (3H, m), 3.90-4.10 (2H, m),
6.40-8.30 (14H, m)

APCI-MASS : m/z = 442 ($\text{M}^+ + 1$)

Preparation 233

1-(E)-[3-[4-[4-(6-Fluorohexyloxy)phenyl]phenyl]-
acryloyl]benzotriazole 3-oxide

IR (KBr) : 3074.0, 3033.5, 2939.0, 2865.7, 1780.0,
1697.1, 1598.7 cm^{-1}

NMR (DMSO- d_6 , δ) : 1.25-1.83 (6H, m), 4.04 (2H, t,
 $J=6.5\text{Hz}$), 4.45 (2H, dt, $J=47.5$ and 6.5Hz), 6.9-8.3
(14H, m)

APCI-MASS : m/z = 460 ($\text{M}^+ + 1$)

Preparation 234

1-(E)-[3-[4-[4-(6-Methoxyhexyloxy)phenyl]phenyl]-
acryloyl]benzotriazole 3-oxide

NMR (DMSO- d_6 , δ) : 1.30-1.65 (6H, m), 1.65-1.90 (2H,
m), 3.22 (3H, s), 3.22-3.40 (2H, m), 4.02 (2H, t,

J=6.5Hz), 6.5-8.3 (14H, m)

Preparation 235

1-[4-[3-(4-n-Hexyloxyphenyl)pyrazol-5-yl]benzoyl]benzotriazole 3-oxide

IR (KBr) : 2935, 1780, 1610, 1506 1249, 1232, 1178,
1087 cm^{-1}

NMR (CDCl_3 , δ) : 0.91 (3H, d, J=6.4Hz), 1.2-1.6 (6H, m), 1.7-1.9 (2H, m), 3.98 (2H, t, J=6.5Hz), 6.8-7.0 (3H, m), 7.4-7.6 (5H, m), 8.00 (2H, d, J=8.4Hz), 8.10 (1H, d, J=8.1Hz), 8.28 (1H, d, J=8.4Hz)

APCI-MASS : m/z = 482 ($\text{M}+\text{H}^+$)

Preparation 236

1-[4-[4-[4-(6-Methoxyhexyloxy)phenyl]phenyl]benzoyl]benzotriazole 3-oxide

IR (KBr) : 2935, 2858, 1774, 1600, 1490, 1257,
1211 cm^{-1}

NMR (CDCl_3 , δ) : 1.4-1.9 (8H, m), 3.35 (3H, s), 3.40 (2H, t, J=6.3Hz), 4.02 (2H, t, J=6.4Hz), 7.00 (2H, d, J=8.7Hz), 7.4-7.8 (7H, m), 7.87 (2H, d, J=8.4Hz), 8.12 (1H, d, J=8.2Hz), 8.36 (2H, d, J=8.4Hz)

APCI-MASS : m/z = 522 ($\text{M}+\text{H}^+$)

Preparation 237

1-[4-[5-[4-(8-Methoxyoctyloxy)phenyl]-1,3,4-thiadiazol-2-yl]benzoyl]benzotriazole 3-oxide

IR (KBr) : 2929, 2854, 1776, 1602, 1469, 1255 cm^{-1}

NMR (CDCl_3 , δ) : 1.2-1.6 (10H, m), 1.7-1.9 (2H, m), 3.33 (3H, s), 3.37 (2H, d, J=6.4Hz), 4.03 (2H, d, J=6.5Hz), 7.00 (2H, d, J=8.9Hz), 7.4-7.6 (3H, m), 7.97 (2H, d, J=8.9Hz), 8.12 (1H, d, J=8.2Hz), 8.23 (2H, d, J=8.7Hz), 8.39 (2H, d, J=8.7Hz)

APCI-MASS : m/z = 558 ($\text{M}+\text{H}^+$)

Preparation 238

1-[4-(4-n-Butoxyphenyl)cinnamoyl]benzotriazole 3-oxide

IR (KBr) : 2952, 2867, 1778, 1598, 1496, 1249,
1186 cm^{-1}

5 NMR (CDCl_3 , δ) : 0.99 (3H, t, $J=7.3\text{Hz}$), 1.55 (2H, tq,
 $J=7.0$ and 7.3Hz), 1.78 (2H, tt, $J=7.0$ and 6.4Hz),
4.02 (2H, t, $J=6.4\text{Hz}$), 6.75 (1H, d, $J=16.0\text{Hz}$), 7.00
(2H, d, $J=8.7\text{Hz}$), 7.4-8.2 (9H, m)

APCI-MASS : $m/z = 414$ ($\text{M}+\text{H}^+$)

10

Preparation 239

1-[4-[5-(4-Cyclohexylphenyl)-1,3,4-thiadiazol-2-yl]benzoyl]benzotriazole 3-oxide

IR (KBr) : 2925, 2850, 1778, 1230, 989 cm^{-1}

15 NMR (CDCl_3 , δ) : 1.2-1.6 (5H, m), 1.7-2.0 (5H, m),
2.5-2.7 (1H, m), 7.37 (2H, d, $J=8.3\text{Hz}$), 7.4-7.6
(3H, m), 7.97 (2H, d, $J=8.3\text{Hz}$), 8.13 (1H, d,
 $J=8.2\text{Hz}$), 8.26 (2H, d, $J=8.6\text{Hz}$), 8.42 (2H, d,
 $J=8.6\text{Hz}$)

20 APCI-MASS : $m/z = 482$ ($\text{M}+\text{H}^+$)

Preparation 240

1-[4-[5-[4-(4-n-Propyloxyphenyl)phenyl]-1,3,4-oxadiazol-2-yl]benzoyl]benzotriazole 3-oxide

25 IR (KBr) : 1778, 1604, 1488, 1249, 1232, 998 cm^{-1}

NMR (CDCl_3 , δ) : 1.07 (3H, t, $J=7.4\text{Hz}$), 1.85 (2H, tq,
 $J=6.5$ and 7.4Hz), 7.02 (2H, d, $J=8.8\text{Hz}$), 7.4-7.7
(3H, m), 7.61 (2H, d, $J=8.8\text{Hz}$), 7.75 (2H, d,
 $J=8.5\text{Hz}$), 8.14 (1H, d, $J=8.2\text{Hz}$), 8.22 (2H, d,
30 $J=8.5\text{Hz}$), 8.40 (2H, d, $J=8.8\text{Hz}$), 8.48 (2H, d,
 $J=8.8\text{Hz}$)

APCI-MASS : $m/z = 518$ ($\text{M}+\text{H}^+$)

Preparation 241

35 1-[4-(5-n-Nonyl-1,3,4-oxadiazol-2-yl)benzoyl]-

benzotriazole 3-oxide

IR (KBr) : 2919, 2850, 1780, 1565, 1415, 1251 cm^{-1}

NMR (CDCl_3 , δ) : 0.89 (3H, t, $J=6.7\text{Hz}$), 1.2-1.6 (12H, m), 1.8-2.0 (2H, m), 2.98 (2H, t, $J=7.7\text{Hz}$), 7.4-7.6 (3H, m), 8.12 (1H, d, $J=9.0\text{Hz}$), 8.28 (2H, d, $J=8.7\text{Hz}$), 8.42 (2H, d, $J=8.7\text{Hz}$)

APCI-MASS : $m/z = 434$ ($\text{M}+\text{H}^+$)

Preparation 242

10 1-[4-[3-(4-n-Hexyloxyphenyl)-1,2,4-oxadiazol-5-yl]-benzoyl]benzotriazole 3-oxide

IR (KBr) : 2946, 2869, 1780, 1251, 1230, 1001 cm^{-1}

15 NMR (CDCl_3 , δ) : 0.92 (3H, t, $J=6.8\text{Hz}$), 1.3-1.6 (6H, m), 1.8-1.9 (2H, m), 4.04 (2H, t, $J=6.5\text{Hz}$), 7.03 (2H, d, $J=8.9\text{Hz}$), 7.4-7.6 (3H, m), 8.0-8.2 (3H, m), 8.46 (4H, s)

APCI-MASS : $m/z = 484$ ($\text{M}+\text{H}^+$)

Preparation 243

20 1-[4-[5-(4-n-Octyloxyphenyl)-1,3,4-thiadiazol-2-yl]-benzoyl]benzotriazole 3-oxide

IR (KBr) : 2925, 2856, 1774, 1602, 1259, 1232, 989 cm^{-1}

25 NMR (CDCl_3 , δ) : 0.90 (3H, t, $J=6.7\text{Hz}$), 1.1-1.6 (10H, m), 1.7-1.9 (2H, m), 4.04 (2H, t, $J=6.5\text{Hz}$), 7.01 (2H, d, $J=8.9\text{Hz}$), 7.4-7.6 (3H, m), 7.97 (2H, d, $J=8.8\text{Hz}$), 8.12 (1H, d, $J=8.2\text{Hz}$), 8.24 (2H, d, $J=8.6\text{Hz}$), 8.40 (2H, d, $J=8.6\text{Hz}$)

APCI-MASS : $m/z = 528$ ($\text{M}+\text{H}^+$)

30 Preparation 244

1-[4-[5-(4-Trans-n-pentylcyclohexyl)-1,3,4-thiadiazol-2-yl]benzoyl]benzotriazole 3-oxide

IR (KBr) : 2952, 2919, 2848, 1785, 1444, 1226, 991 cm^{-1}

35 NMR (CDCl_3 , δ) : 0.90 (3H, t, $J=6.9\text{Hz}$), 1.0-1.7 (13H, m), 1.94 (2H, d, $J=12.0\text{Hz}$), 2.27 (2H, d, $J=12.0\text{Hz}$),

5

Preparation 245

10

NMR (CDCl₃, δ) : 0.95 (3H, t, J=7.1Hz), 1.2-1.6 (4H, m), 1.7-1.9 (2H, m), 4.02 (2H, t, J=6.5Hz), 7.0-7.1 (3H, m), 7.4-7.6 (3H, m), 7.81 (2H, d, J=8.8Hz), 8.06 (2H, d, J=8.6Hz), 8.12 (1H, d, J=8.0Hz), 8.39 (2H, d, J=8.6Hz)

15

Preparation 246

20

NMR (CDCl₃, δ) : 1.2-1.6 (10H, m), 1.7-1.9 (2H, m), 3.34 (3H, s), 3.38 (2H, t, J=6.4Hz), 4.05 (2H, t, J=6.5Hz), 7.04 (2H, d, J=8.8Hz), 7.4-7.6 (3H, s), 8.1-8.2 (3H, s), 8.36 (2H, d, J=8.7Hz), 8.45 (2H, d, J=8.7Hz)

25

Preparation 247

30

IR (KBr) : 1783, 1604, 1423, 1284, 985 cm^{-1}

NMR (CDCl₃, δ) : 7.2-8.2 (15H, m), 8.12 (2H, d, J=8.3Hz), 8.36 (2H, d, J=8.4Hz)

35

APCI-MASS : $m/z = 486$ ($M^+ + 1$)

Preparation 248

1-[4-[5-(4-n-Octyloxyphenyl)-1,3,4-oxadiazol-2-yl]benzoyl]benzotriazole 3-oxide

IR (KBr) : 2925, 2854, 1780, 1610, 1496, 1257, 1228,
1180 cm^{-1}

NMR (CDCl_3 , δ) : 0.89 (3H, t, $J=6.8\text{Hz}$), 1.2-2.0 (12H, m), 4.05 (2H, t, $J=6.5\text{Hz}$), 7.05 (2H, d, $J=8.7\text{Hz}$), 7.4-7.6 (3H, m), 8.0-8.2 (3H, m), 8.37 (2H, d, $J=8.6\text{Hz}$), 8.45 (2H, d, $J=8.6\text{Hz}$)

APCI-MASS : $m/z = 512$ ($\text{M}+\text{H}^+$)

Preparation 249

1-[4-[2-(4-n-Hexyloxyphenyl)pyrimidin-6-yl]benzoyl]benzotriazole 3-oxide

IR (KBr) : 2948, 2861, 1780, 1552, 1413, 1378, 987 cm^{-1}

NMR (CDCl_3 , δ) : 0.92 (3H, t, $J=6.8\text{Hz}$), 1.2-1.6 (6H, m), 1.8-2.0 (2H, m), 4.06 (2H, t, $J=6.5\text{Hz}$), 7.04 (2H, d, $J=9.0\text{Hz}$), 7.4-7.6 (3H, m), 7.64 (1H, d, $J=5.2\text{Hz}$), 8.13 (1H, d, $J=8.2\text{Hz}$), 8.44 (4H, s), 8.55 (2H, d, $J=9.0\text{Hz}$), 8.90 (1H, d, $J=5.2\text{Hz}$)

APCI-MASS : $m/z = 494$ ($\text{M}+\text{H}^+$)

Preparation 250

1-[4-[4-[8-(2-Ethoxyethoxy)octyloxy]phenyl]benzoyl]benzotriazole 3-oxide

IR (KBr) : 2933, 2861, 1778, 1598, 1247, 1186, 977 cm^{-1}

NMR (CDCl_3 , δ) : 1.22 (3H, t, $J=7.0\text{Hz}$), 1.3-2.0 (14H, m), 3.4-3.6 (6H, m), 4.02 (2H, t, $J=6.5\text{Hz}$), 7.02 (2H, d, $J=8.8\text{Hz}$), 7.4-7.6 (3H, m), 7.62 (2H, d, $J=8.8\text{Hz}$), 7.78 (2H, d, $J=8.6\text{Hz}$), 8.10 (1H, d, $J=8.9\text{Hz}$), 8.31 (2H, d, $J=8.6\text{Hz}$)

APCI-MASS : $m/z = 532$ ($\text{M}+\text{H}^+$)

Preparation 251

1-[4-[4-[7-(Piperidin-1-yl-carbonyl)heptyloxy]phenyl]-

benzoyl]benzotriazole 3-oxide

IR (KBr) : 2935, 2856, 1774, 1631, 1598, 1255,
1191 cm^{-1}

5 NMR (CDCl_3 , δ) : 1.3-2.0 (16H, m), 2.37 (2H, t,
J=7.6Hz), 3.48 (4H, s), 4.02 (2H, t, J=6.4Hz), 7.02
(2H, d, J=8.6Hz), 7.4-7.6 (3H, m), 7.63 (2H, d,
J=8.6Hz), 7.78 (2H, d, J=8.3Hz), 8.11 (1H, d,
J=8.1Hz), 8.31 (2H, d, J=8.3Hz)

10 APCI-MASS : m/z = 541 ($\text{M}+\text{H}^+$)

Preparation 252

1-[6-[4-(4-n-Heptyloxyphenyl)piperazin-1-yl]nicotinoyl]-
benzotriazole 3-oxide

IR (KBr) : 2929, 2856, 1762, 1604, 1510, 1240 cm^{-1}

15 NMR (CDCl_3 , δ) : 0.89 (3H, t, J=6.7Hz), 1.2-1.9 (10H,
m), 3.20 (4H, t, J=5.0Hz), 3.8-4.0 (6H, m), 6.75
(1H, d, J=9.5Hz), 6.86 (2H, d, J=9.3Hz), 6.95 (2H,
d, J=9.3Hz), 7.3-7.6 (3H, m), 8.10 (1H, d,
J=8.2Hz), 8.19 (1H, dd, J=9.2 and 2.3Hz), 9.05 (1H,
20 d, J=2.3Hz)

APCI-MASS : m/z = 515 ($\text{M}+\text{H}^+$)

Preparation 253

25 1-[6-[4-[4-(8-Methoxyoctyloxy)phenyl]piperazin-1-
yl]nicotinoyl]benzotriazole 3-oxide

IR (KBr) : 2929, 2854, 1766, 1602, 1510, 1419,
1234 cm^{-1}

30 NMR (CDCl_3 , δ) : 1.3-1.9 (12H, m), 3.2-3.3 (4H, m),
3.33 (3H, s), 3.36 (2H, t, J=6.4Hz), 3.92 (2H, t,
J=6.5Hz), 4.0-4.2 (4H, m), 6.75 (1H, d, J=9.1Hz),
6.87 (2H, d, J=8.9Hz), 7.0-7.2 (2H, m), 7.4-7.6
(3H, m), 8.09 (1H, d, J=8.1Hz), 8.20 (1H, dd, J=9.1
and 2.3Hz), 9.05 (1H, d, J=2.3Hz)

35 APCI-MASS : m/z = 559 ($\text{M}+\text{H}^+$)

Preparation 254

1-[4-[5-[4-(4-n-Propyloxyphenyl)phenyl]-1,3,4-thiadiazol-2-yl]benzoyl]benzotriazole 3-oxide

IR (KBr) : 1774, 1600, 1234, 985 cm^{-1}

5 NMR (CDCl_3 , δ) : 1.07 (3H, t, $J=7.3\text{Hz}$), 1.85 (2H, tq, $J=6.5$ and 7.3Hz), 3.99 (2H, t, $J=6.5\text{Hz}$), 7.01 (2H, d, $J=8.7\text{Hz}$), 7.4-7.7 (5H, m), 7.72 (2H, d, $J=8.7\text{Hz}$), 8.1-8.2 (2H, m), 8.28 (2H, d, $J=8.6\text{Hz}$), 8.44 (2H, d, $J=8.6\text{Hz}$)

10 APCI-MASS : $m/z = 534$ ($M+H$)⁺

The following compounds (Preparations 255 to 256) were obtained according to a similar manner to that of Preparation 32.

15

Preparation 255

6-Heptylnaphthalene-2-carboxylic acid

20 NMR (CDCl_3 , δ) : 0.88 (3H, t, $J=6.6\text{Hz}$), 1.15-1.53 (8H, m), 1.58-1.88 (2H, m), 2.80 (2H, t, $J=7.6\text{Hz}$), 7.42 (1H, dd, $J=1.7$ and 8.4Hz), 7.67 (1H, s), 7.84 (1H, d, $J=8.6\text{Hz}$), 7.90 (1H, d, $J=8.4\text{Hz}$), 8.09 (1H, dd, $J=1.7$ and 8.6Hz), 8.68 (1H, s)

APCI-MASS : $m/z = 271$ (M^++1), 285 (methyl ester⁺-1)

25

Preparation 256

3-(E)-[4-[4-(7-Fluoroheptyloxy)phenyl]phenyl]acrylic acid

IR (KBr) : 3037.3, 2935.1, 2861.8, 1679.7, 1633.4, 1600.6 cm^{-1}

30

NMR ($\text{DMSO}-d_6$, δ) : 1.30-1.85 (10H, m), 4.01 (2H, t, $J=6.4\text{Hz}$), 4.44 (2H, dt, $J=47.6$ and 6.1Hz), 6.54 (1H, d, $J=15.9\text{Hz}$), 7.02 (2H, d, $J=8.7\text{Hz}$), 7.53-7.80 (7H, m)

35

Preparation 257

To a solution of 4-methylpentanol (3.0 ml) in pyridine (20 ml) were added in turn with p-toluenesulfonyl chloride (4.6 g) and 4-N,N-dimethylaminopyridine (1.5 g) at ambient temperature. After stirring at ambient temperature, the reaction mixture was taken up into a mixture of ethyl acetate (100 ml) and water (100 ml). The separated organic layer was washed in turn with hydrochloric acid(1N), water, aqueous sodium hydrogencarbonate, and brine, and dried over magnesium sulfate. Evaporation gave 1-p-Toluenesulfonyloxy-4-methylpentane (5.30 g).

NMR (CDCl₃, δ) : 0.83 (6H, d, J=6.6Hz), 1.48 (1H, sept, J=6.6Hz), 1.50-1.70 (2H, m), 2.45 (3H, s), 4.00 (2H, t, J=6.6Hz), 7.34 (2H, d, J=8.1Hz), 7.79 (2H, d, J=8.1Hz)

APCI-MASS : m/z = 257 (M⁺+1)

Preparation 258

To a solution of 4-bromo-4'-n-butyloxybiphenyl (3.05 g) in tetrahydrofuran (60 ml) was added 1.55M n-butyllithium in n-hexane (7.74 ml) at -60°C over a period of 10 minutes. The solution was stirred at -30°C for 1.5 hours and cooled to -60°C. To the solution was added triisopropylborate (3.46 ml) over a period of 5 minutes, and the mixture was stirred for 1.5 hours without cooling. To the solution was added 1N hydrochloric acid (20 ml) and the solution was stirred for 30 minutes and extracted with ethyl acetate. The organic layer was separated and washed with water, brine and dried over magnesium sulfate. The solvents were removed under reduced pressure and the residue was triturated with n-hexane. The solid was collected by filtration and dried under reduced pressure to give 4-(4-n-Butyloxyphenyl)phenylboronic acid (2.31 g).

IR (KBr) : 3398, 2956, 2919, 2871, 1604, 1531, 1392, 1257 cm⁻¹

NMR (DMSO-d₆, δ) : 0.94 (3H, t, J=7.3Hz), 1.4-1.8 (4H,

m), 4.01 (2H, t, J=6.3Hz), 7.01 (2H, d, J=8.7Hz),
7.58 (2H, d, J=7.9Hz), 7.62 (2H, d, J=8.7Hz), 7.84
(2H, d, J=7.9Hz), 8.03 (2H, s)

5 The following compounds (Preparations 259 to 260) were
obtained according to a similar manner to that of Preparation
258.

Preparation 259

10 4-[4-(6-Methoxyhexyloxy)phenyl]phenylboronic acid
IR (KBr) : 3448, 3392, 2937, 2861, 1606, 1529, 1346,
1288 cm^{-1}
NMR (DMSO- d_6 , δ) : 1.3-1.8 (8H, m), 3.21 (3H, s), 3.31
(2H, t, J=6.3Hz), 3.99 (2H, t, J=6.4Hz), 7.00 (2H,
15 d, J=8.7Hz), 7.5-7.7 (4H, m), 7.84 (2H, d,
J=8.1Hz), 8.03 (2H, s)
APCI-MASS : m/z = 329 ($\text{M}+\text{H}^+$)

Preparation 260

20 4-[4-(5-Methoxypentyloxy)phenyl]phenylboronic acid
IR (KBr) : 3473, 3369, 3330, 2935, 2863, 1604, 1531,
1338, 1251 cm^{-1}
NMR (DMSO- d_6 , δ) : 1.4-1.8 (6H, m), 3.22 (3H, s), 3.3-
3.4 (2H, m), 3.99 (2H, t, J=6.4Hz), 7.00 (2H, d,
25 J=8.7Hz), 7.58 (2H, d, J=8.0Hz), 7.61 (2H, d,
J=8.7Hz), 7.84 (2H, d, J=8.0Hz), 8.04 (2H, s)
APCI-MASS : m/z = 315 ($\text{M}+\text{H}^+$)

Preparation 261

30 To a suspension of 4-Methoxycarbonylphenyl boronic acid
(648 mg) and 4-iodo-1-heptylpyrazole (876 mg) and $\text{Pd}(\text{PPh}_3)_4$
(173 mg) in 1,2-dimethoxyethane (10 ml) was added 2M Na_2CO_3
aq. (3.6 ml). The reaction mixture was stirred at 80°C for 2
hours under N_2 atmosphere, and poured into ice-water and
35 extracted with ethyl acetate. The organic layer was washed

with brine, and dried over MgSO_4 . The solvent was removed under pressure. The residue was subjected to column-chromatography on silica gel 60 (Merk) and eluted with n-hexane/ethyl acetate (80:20). The fractions containing the object compound were combined and evaporated under reduced pressure to give 1-heptyl-4-(4-methoxycarbonylphenyl)pyrazole (0.20 g).

IR (KBr pelet) : 2952, 2920, 2848, 1712, 1610, 1288, 1114, 769 cm^{-1}

NMR (DMSO-d_6 , δ) : 0.85 (3H, t, $J=6.7\text{Hz}$), 1.1-1.4 (8H, m), 1.7-1.9 (2H, m), 3.85 (3H, s), 4.11 (2H, t, $J=7.0\text{Hz}$), 7.72 (2H, d, $J=8.5\text{Hz}$), 7.93 (2H, d, $J=8.5\text{Hz}$), 7.99 (1H, s), 8.34 (1H, s)

APCI-MASS : $m/z = 301$ ($\text{M}+\text{H}^+$)

The following compounds (Preparations 262 to 268) were obtained according to a similar manner to that of Preparation 261.

Preparation 262

Ethyl 4-[4-(4-n-butyloxyphenyl)phenyl]benzoate

IR (KBr) : 2958, 2935, 2871, 1714, 1602, 1396, 1280, 1108 cm^{-1}

NMR (CDCl_3 , δ) : 0.99 (3H, t, $J=7.3\text{Hz}$), 1.4-2.0 (7H, m), 4.02 (2H, t, $J=6.4\text{Hz}$), 4.40 (2H, q, $J=7.1\text{Hz}$), 6.98 (2H, d, $J=6.8\text{Hz}$), 7.56 (2H, d, $J=6.8\text{Hz}$), 7.66 (4H, s), 7.68 (2H, d, $J=8.4\text{Hz}$), 8.12 (2H, d, $J=8.4\text{Hz}$)

APCI-MASS : $m/z = 375$ ($\text{M}+\text{H}^+$)

Preparation 263

Methyl 6-(4-heptyloxyphenyl)nicotinate

IR (KBr) : 2954, 2859, 1724, 1597, 1288, 1251, 1116, 783 cm^{-1}

NMR (CDCl_3 , δ) : 0.90 (3H, t, $J=6.6\text{Hz}$), 1.2-1.5 (8H,

m), 1.7-1.9 (2H, m), 3.96 (3H, s), 4.03 (2H, t, J=6.5Hz), 7.00 (2H, d, J=8.8Hz), 7.75 (1H, d, J=8.4Hz), 8.02 (1H, d, J=8.8Hz), 8.30 (1H, dd, J=8.4 and 2.2Hz), 9.23 (1H, d, J=2.2Hz)

5 APCI-MASS : m/z = 328 (M+H⁺)

Preparation 264

Methyl 6-[4-(4-n-butyloxyphenyl)phenyl]nicotinate

10 IR (KBr) : 2956, 2933, 2871, 1724, 1598, 1282,
1118 cm⁻¹

NMR (CDCl₃, δ) : 1.00 (3H, t, J=7.3Hz), 1.4-1.9 (4H, m), 3.98 (3H, s), 4.02 (2H, t, J=6.4Hz), 7.00 (2H, d, J=8.8Hz), 7.59 (2H, d, J=8.8Hz), 7.70 (2H, d, J=8.5Hz), 7.86 (1H, d, J=8.8Hz), 8.13 (2H, d, J=8.5Hz), 8.37 (1H, dd, J=8.8 and 1.6Hz), 9.30 (1H, d, J=1.6Hz)

15 APCI-MASS : m/z = 362 (M+H⁺)

Preparation 265

20 Methyl 5-[4-(4-n-butyloxyphenyl)phenyl]furan 2-carboxylate

IR (KBr) : 2958, 2933, 2873, 1716, 1483, 1303,
1139 cm⁻¹

25 NMR (CDCl₃, δ) : 0.99 (3H, t, J=7.3Hz), 1.5-1.9 (4H, m), 3.93 (3H, s), 4.01 (2H, t, J=6.4Hz), 6.75 (1H, d, J=3.6Hz), 6.98 (2H, d, J=8.7Hz), 7.26 (1H, d, J=3.6Hz), 7.56 (2H, d, J=8.4Hz), 7.61 (2H, d, J=8.7Hz), 7.83 (2H, d, J=8.4Hz)

30 APCI-MASS : m/z = 351 (M+H)⁺

Preparation 266

Ethyl 4-[4-[4-(6-methoxyhexyloxy)phenyl]phenyl]benzoate

IR (KBr) : 2937, 2863, 1712, 1602, 1396, 1278,
1108 cm⁻¹

35 NMR (CDCl₃, δ) : 1.4-2.0 (11H, m), 3.34 (3H, s), 3.39

(2H, t, J=6.4Hz), 4.01 (2H, t, J=6.4Hz), 4.41 (2H, q, J=7.1Hz), 6.98 (2H, d, J=8.7Hz), 7.56 (2H, d, J=8.7Hz), 7.6-7.8 (6H, m), 8.12 (2H, d, J=8.4Hz)
APCI-MASS : m/z = 433 (M+H⁺)

5

Preparation 267

4-[4-[4-(5-Methoxypentyloxy)phenyl]phenyl]benzoic acid

IR (KBr) : 2939, 2859, 1679, 1587, 1396, 1321, 1292, 1126 cm⁻¹

10

NMR (DMSO-d₆, δ) : 1.3-1.8 (6H, m), 3.21 (3H, s), 3.2-3.4 (2H, m), 4.01 (2H, t, J=6.5Hz), 7.04 (2H, d, J=8.6Hz), 7.66 (2H, d, J=8.6Hz), 7.7-7.9 (6H, m), 8.03 (2H, d, J=8.2Hz)

APCI-MASS : m/z = 391 (M+H⁺)

15

Preparation 268

Methyl 4-[4-[4-(5-methoxypentyloxy)phenyl]phenyl]phenyl acetate

IR (KBr) : 2937, 2863, 1739, 1604, 1492, 1255 cm⁻¹

20

NMR (CDCl₃, δ) : 1.5-2.0 (6H, m), 3.34 (3H, s), 3.42 (2H, t, J=6.3Hz), 3.68 (2H, s), 3.72 (3H, s), 4.02 (2H, t, J=6.4Hz), 6.97 (2H, d, J=8.7Hz), 7.36 (2H, d, J=8.2Hz), 7.5-7.7 (8H, m)

APCI-MASS : m/z = 419 (M+H⁺)

25

Preparation 269

A solution of 3-[2-(4-Hexylphenylamino)ethyl]-2-oxo-oxazolidine hydrochloride (2.131 g) in 25% hydrobromic acid in acetic acid (13.04 ml) was stirred for 96 hours at ambient temperature. The reaction mixture was pulverized with diisopropyl ether. The precipitate was collected by filtration and added to ethanol (15 ml). The solution was refluxed for 5 hours and pulverized with diisopropyl ether. The precipitate was collected by filtration to give 1-(4-n-Hexylphenyl)piperazine dihydrobromide (2.413 g).

35

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IR (KBr) : 2921.6, 2711.4, 2485.8, 1452.1, 1012.4 cm^{-1}
NMR (DMSO- d_6 , δ) : 0.85 (3H, t, $J=6.6\text{Hz}$), 1.1-1.4 (6H, m), 1.4-1.6 (2H, m), 2.49 (2H, t, $J=8.4\text{Hz}$), 3.1-3.4 (8H, m), 6.54 (2H, s), 6.90 (2H, d, $J=8.7\text{Hz}$), 7.08 (2H, d, $J=8.7\text{Hz}$), 8.78 (1H, s)
APCI-MASS : $m/z = 247$ ($\text{M}^+ + \text{H}$)

The following compounds (Preparations 270 to 274) were obtained according to a similar manner to that of Preparation 269.

Preparation 270

4-[4-(4-n-Hexylphenyl)piperazin-1-yl]benzoic acid dihydrobromide

IR (KBr) : 2956.3, 1691.3, 1664.3, 1602.6, 1232.3 cm^{-1}
NMR (DMSO- d_6 , δ) : 0.85 (3H, t, $J=6.5\text{Hz}$), 1.2-1.4 (10H, m), 1.4-1.6 (2H, m), 2.51 (2H, t, $J=7.4\text{Hz}$), 3.2-3.6 (8H, m), 7.0-7.2 (6H, m), 7.81 (2H, d, $J=8.8\text{Hz}$)

APCI-MASS : $m/z = 367$ ($\text{M}^+ + \text{H}$)

Preparation 271

1-(4-Cyclohexylphenyl)piperazine dihydrobromide

IR (KBr) : 2927.4, 1510.0, 1452.1 cm^{-1}
NMR (DMSO- d_6 , δ) : 1.1-1.5 (6H, m), 1.6-1.9 (4H, m), 2.41 (1H, m), 3.1-3.4 (8H, m), 6.91 (2H, d, $J=8.7\text{Hz}$), 7.11 (2H, d, $J=8.7\text{Hz}$), 8.78 (1H, s)
APCI-MASS : $m/z = 245$ ($\text{M}^+ + \text{H}$)

Preparation 272

4-[4-(4-Cyclohexylphenyl)piperazin-1-yl]benzoic acid dihydrobromide

IR (KBr) : 1668.1, 1602.6, 1230.4, 1189.9 cm^{-1}
APCI-MASS : $m/z = 365$ ($\text{M}^+ + \text{H}$)

Preparation 273

3-Fluoro-4-[4-(4-hydroxyphenyl)piperazin-1-yl]benzoic acid dihydrobromide

IR (KBr) : 1708.6, 1610.3 cm^{-1}

5 NMR (DMSO-d_6 , δ) : 3.2-3.6 (8H, m), 6.81 (2H, d, $J=8.6\text{Hz}$), 7.0-7.4 (3H, m), 7.4-7.8 (2H, m)

APCI-MASS : $m/z = 317$ ($M^+ + H$)

Preparation 274

10 4-[4-(4-Hydroxyphenyl)piperazin-1-yl]benzoic acid dihydrobromide

IR (KBr) : 1670.1, 1604.5, 1226.5, 775.2 cm^{-1}

15 NMR (DMSO-d_6 , δ) : 3.0-3.2 (4H, m), 3.3-3.5 (4H, m), 6.68 (2H, d, $J=8.8\text{Hz}$), 6.85 (2H, d, $J=8.8\text{Hz}$), 7.02 (2H, d, $J=8.8\text{Hz}$), 7.79 (2H, d, $J=8.8\text{Hz}$), 8.86 (1H, s), 12.29 (1H, s)

APCI-MASS : $m/z = 299$ ($M + H^+$)

Preparation 275

20 A mixture of 4-n-hexyloxyaniline (10 g), ethyl acrylate (56.1 ml), glacial acetic acid (19.25 ml), and cuprous chloride (1.02 g) was heated under reflux with stirring under nitrogen for 26 hours. A solution of the cold product in ether was shaken with water and then with aqueous ammonia.
25 The organic layer was taken and dried over magnesium sulfate. The magnesium sulfate was filtered off, and filtrate was evaporated under reduced pressure. The residue was subjected to column chromatography on silica gel and eluted with hexane - ethyl acetate (9:1). The fractions containing the
30 object compound were combined and evaporated under reduced pressure to give Ethyl 3-[N-(2-ethoxycarbonyl)ethyl]-N-(4-hexyloxyphenyl)amino]propionate (15.756 g).

IR (Neat) : 1733.7, 1513.8, 1241.9, 1182.2 cm^{-1}

35 NMR (CDCl_3 , δ) : 0.90 (3H, t, $J=6.5\text{Hz}$), 1.2-1.55 (6H, m), 1.24 (6H, t, $J=7.1\text{Hz}$), 1.65-1.85 (2H, m), 2.51

(4H, t, J=7.2Hz), 3.53 (4H, t, J=7.2Hz), 3.89 (2H, t, J=6.5Hz), 4.12 (4H, q, J=7.1Hz), 6.72 (2H, d, J=9.3Hz), 6.83 (2H, d, J=9.3Hz)

APCI-MASS : $m/z = 394 (M^+ + H)$

5

Preparation 276

A suspension of methyl 4-formylbenzoate (4.92 g), hydroxylamine hydrochloride (5.21 g) and sodium acetate (6.15 g) in ethanol (50 ml) was refluxed for 2 hours. The mixture was poured into water and extracted with ethyl acetate and the separated organic layer was washed with brine and dried over magnesium sulfate. The solvents were removed under reduced pressure to give 4-methoxycarbonylbenzaldehyde oxime (5.28 g).

10

15

IR (KBr) : 3291, 1727, 1438, 1284, 1112 cm^{-1}

NMR (CDCl_3 , δ) : 3.93 (3H, s), 7.65 (2H, d, J=8.3Hz), 8.10 (2H, d, J=8.3Hz), 8.18 (1H, s), 8.27 (1H, s)

APCI-MASS : $m/z = 180$

20

The following compound was obtained according to a similar manner to that of Preparation 276.

Preparation 277

N-Hydroxy-4-n-hexyloxybenzamidinium

25

IR (KBr) : 3446, 3349, 2937, 2865, 1650, 1610, 1519, 1392, 1253 cm^{-1}

NMR ($\text{DMSO}-d_6$, δ) : 0.88 (3H, t, J=6.4Hz), 1.2-1.8 (8H, m), 3.97 (2H, t, J=6.5Hz), 5.70 (2H, s), 6.90 (2H, d, J=8.4Hz), 7.58 (2H, d, J=8.4Hz), 9.43 (1H, s)

30

APCI-MASS : $m/z = 237 (M+H)^+$

Preparation 278

To a solution of 4-methoxycarbonylbenzaldehyde oxime (896 mg) in N,N-dimethylformamide (10 ml) was added 4N-hydrochloride acid in 1,4-dioxane (1.38 ml) and oxone^R (1.69

35

g). The suspension was stirred at ambient temperature for 16 hours and poured into ice-water. The object compound was extracted with ethyl acetate and the organic layer was washed with brine, dried over magnesium sulfate. The solvents were removed under reduced pressure to give 4-Methoxycarbonylbenzaldehyde oxime chloride (1.05 g).

IR (KBr) : 3390, 1710, 1436, 1405, 1284, 1232, 1116, 1016 cm^{-1}

NMR (CDCl_3 , δ) : 3.95 (3H, s), 8.93 (2H, d, $J=8.3\text{Hz}$), 8.10 (2H, d, $J=8.7\text{Hz}$), 8.39 (1H, s)

APCI-MASS : $m/z = 176$ ($\text{M-H}^+-\text{HCl}$)

Preparation 279

A solution of Ethyl 4-oxo-1-(4-n-hexyloxyphenyl)piperidine-3-carboxylate (1.437 g) in 20% hydrochloric acid (7.2 ml) was refluxed for 2 hours, cooled, basified with 60% aqueous sodium hydroxide, and extracted with ethyl acetate. The organic layer was taken and dried over magnesium sulfate. The magnesium sulfate was filtered off, and filtrate was evaporated under reduced pressure to give 1-(4-n-Hexyloxyphenyl)-4-piperidone (0.959 g).

IR (Neat) : 2931.3, 1716.3, 1511.9, 1243.9, 825.4 cm^{-1}

NMR (CDCl_3 , δ) : 0.90 (3H, t, $J=6.5\text{Hz}$), 1.2-1.6 (6H, m), 1.65-1.85 (2H, m), 2.57 (4H, t, $J=6.1\text{Hz}$), 3.46 (4H, t, $J=6.1\text{Hz}$), 3.92 (2H, t, $J=6.5\text{Hz}$), 6.85 (2H, d, $J=9.3\text{Hz}$), 6.95 (2H, d, $J=9.3\text{Hz}$)

APCI-MASS : $m/z = 276$ (M^++H)

Preparation 280

A solution of 4-[4-(7-Bromoheptyloxy)phenyl]bromobenzene (0.25 g) in a solution of tetra n-butylammonium fluoride (tetrahydrofuran solution, 1M, 2.9 ml) was heated to 50°C for 2 hours. After cooling to ambient temperature, the solution was taken up into a mixture of ethyl acetate (20 ml) and water (20 ml). The separated organic layer was washed with

water, brine, and dried over magnesium sulfate. Evaporation gave a residue which was chromatographed on silica gel (30 ml) eluting with a mixture of n-hexane and ethyl acetate (100:0-97:3, V/V). The fractions which contained the objective compound were collected and evaporated a residue which was triturated with n-hexane to give 4-[4-(7-Fluoroheptyloxy)phenyl]bromobenzene (104 mg).

IR (KBr) : 2937.1, 2859.9, 1606.4 cm^{-1}

NMR (CDCl_3 , δ) : 1.20-1.90 (10H, m), 3.99 (2H, t, $J=6.4\text{Hz}$), 4.45 (2H, dt, $J=47.3$ and 6.1Hz), 6.95 (2H, d, $J=6.7\text{Hz}$), 7.40 (2H, d, $J=6.7\text{Hz}$), 7.47 (2H, d, $J=6.7\text{Hz}$), 7.52 (2H, d, $J=6.7\text{Hz}$)

The following compound was obtained according to a similar manner to that of Preparation 280.

Preparation 281

4-[4-(6-Fluorohexyloxy)phenyl]bromobenzene

NMR (CDCl_3 , δ) : 1.40-1.95 (8H, m), 4.01 (2H, t, $J=6.4\text{Hz}$), 4.47 (2H, dt, $J=47.5$ and 6.0Hz), 6.95 (2H, d, $J=8.6\text{Hz}$), 7.35-7.59 (6H, m)

Preparation 282

A solution of 4-[4-(8-Bromooctyloxy)phenyl]bromobenzene (3.7 g) in a mixture of sodium methoxide (4.9M in methanol, 17 ml), N,N-dimethylformamide (20 ml) and tetrahydrofuran (8 ml) was heated to 80°C for 3 hours. The reaction mixture was taken up into a mixture of ethyl acetate (200 ml) and water (100 ml). The separated organic layer was washed in turn with water, brine, dried over magnesium sulfate. Evaporation gave a residue which was subjected to column chromatography (silica gel, 100 ml) eluting with n-hexane to give 4-[4-(8-Methoxyoctyloxy)phenyl]bromobenzene (2.73 g).

IR (KBr) : 2935.1, 2858.0, 1604.5 cm^{-1}

NMR (CDCl_3 , δ) : 1.25-1.70 (10H, m), 1.70-1.95 (2H,

m), 3.33 (3H, s), 3.37 (2H, t, J=6.5Hz), 3.99 (2H, t, J=6.5Hz), 6.95 (2H, d, J=8.8Hz), 7.35-7.66 (6H, m)

APCI-MASS : m/z = 391 (M⁺)

5

The following compounds (Preparations 283 to 284) were obtained according to a similar manner to that of Preparation 282.

10 Preparation 283

4-[4-(6-Methoxyhexyloxy)phenyl]bromobenzene

NMR (CDCl₃, δ) : 1.50-1.70 (6H, m), 1.70-1.95 (2H, m), 3.34 (3H, s), 3.40 (2H, t, J=6.2Hz), 3.99 (2H, t, J=6.5Hz), 6.95 (2H, d, J=8.7Hz), 7.30-7.60 (6H, m)

15 APCI-MASS : m/z = 365 (M⁺+2)

Preparation 284

4-[4-(7-Methoxyheptyloxy)phenyl]bromobenzene

IR (KBr) : 2935.1, 2854.1, 1604.5 cm⁻¹

20 NMR (CDCl₃, δ) : 1.25-1.70 (8H, m), 1.70-1.95 (2H, m), 3.33 (3H, s), 3.37 (2H, t, J=6.4Hz), 3.98 (2H, t, J=6.5Hz), 6.95 (2H, d, J=8.8Hz), 7.35-7.56 (6H, m)

APCI-MASS : m/z = 379 (M⁺+2)

25 Preparation 285

N-(4-octylphenyl)-N'-aminourea, Formamidine acetate (12.76 g) and N-carbazoyl-4-octylaniline (6.458 g) in N,N-dimethylformamide (19.4 ml) were stirred at 150°C for 6 hours. The reaction mixture was pulverized with water. The precipitate was collected by filtration and washed with water to give 4-(4-Octylphenyl)-2,3-dihydro-4H-1,2,4-triazol-3-one (4.27 g).

IR (KBr) : 3214.8, 3085.5, 1704.8 cm⁻¹

35 NMR (CDCl₃, δ) : 0.88 (3H, t, J=6.7Hz), 1.2-1.5 (10H, m), 1.5-1.8 (2H, m), 2.64 (2H, t, J=7.9Hz), 7.29

APCI-MASS : $m/z = 274$ ($M+H^+$)

5

Preparation 286

10

IR (KBr) : 3200, 1699.0, 918.0 cm^{-1}

NMR (CDCl₃, δ) : 1.49 (9H, s), 3.17 (4H, t, J=4.9Hz), 3.60 (4H, t, J=4.9Hz), 7.00 (2H, d, J=9.0Hz), 7.40 (2H, d, J=9.0Hz), 7.63 (1H, s), 10.4 (1H, s)

15

Preparation 287

20

NMR (CDCl₃, δ) : 0.88 (3H, t, J=6.6Hz), 1.16-1.50 (8H, m), 1.50-1.80 (2H, m), 2.78 (2H, t, J=7.6Hz), 3.97 (3H, s), 7.39 (1H, dd, J=17 and 8.4Hz), 7.64 (1H, s), 7.79 (1H, d, J=8.6Hz), 7.86 (1H, d, J=8.4Hz), 8.02 (1H, dd, J=1.7 and 8.6Hz), 8.57 (1H, s)

APCI-MASS : $m/z = 285$ ($M^+ + 1$)

30

The following compound (Preparation 288) was obtained according to a similar manner to that of Preparation 287.

Preparation 288

35

NMR (CDCl_3 , δ) : 0.88 (3H, t, $J=6.8\text{Hz}$), 1.17-1.53 (6H, m), 1.60-1.82 (2H, m), 2.79 (2H, t, $J=7.7\text{Hz}$), 3.97 (3H, s), 7.39 (1H, dd, $J=1.7$ and 8.4Hz), 7.64 (1H, s), 7.80 (1H, d, $J=8.6\text{Hz}$), 7.86 (1H, d, $J=8.4\text{Hz}$), 8.03 (1H, dd, $J=1.7$ and 8.6Hz), 8.57 (1H, s)

APCI-MASS : $m/z = 271$ ($M+1$)

Preparation 289

To a stirred solution of Methyl 6-hydroxynaphthalene-2-carboxylate (3.0 g) in dichloromethane (40 ml) were added in turn diisopropylethylamine (3.9 ml) and triflic anhydride (3.0 ml) at -40°C . After stirring at -40°C for 20 minutes, the mixture was taken up into a mixture of ethyl acetate and cold water. The organic layer was separated, washed with brine, dried over magnesium sulfate, and dried in vacuo. The residue was taken up into piperidine (20 ml) and to the solution were added 1-heptyne (4.0 ml) and tetrakis(triphenylphosphine)palladium(0) (0.5 g). After heating to 85°C for 1 hour under nitrogen atmosphere, the reaction mixture was evaporated in vacuo. The residue was diluted with ethyl acetate, and the solution was washed in turn with hydrochloric acid and brine, dried over magnesium sulfate and evaporated in vacuo. The residue was chromatographed on silica gel (200 ml) eluting with a mixture of n-hexane and ethyl acetate (9:1, V/V) to give Methyl 6-(1-heptynyl)naphthalene-2-carboxylate (4.01 g).

NMR (CDCl_3 , δ) : 0.94 (3H, t, $J=7.1\text{Hz}$), 1.30-1.70 (6H, m), 2.46 (2H, t, $J=7.0\text{Hz}$), 3.97 (3H, s), 7.50 (1H, dd, $J=1.7$ and 8.6Hz), 7.80 (1H, d, $J=8.6\text{Hz}$), 7.86 (1H, d, $J=8.6\text{Hz}$), 8.04 (1H, dd, $J=1.7$ and 8.6Hz), 8.55 (1H, s)

APCI-MASS : $m/z = 281$ (M^++1)

The following compound was obtained according to a similar manner to that of Preparation 289.

Preparation 290

Methyl 6-(1-hexynyl)naphthalene-2-carboxylate

NMR (CDCl₃, δ) : 0.97 (3H, t, J=7.1Hz), 1.40-1.71 (4H, m), 2.47 (2H, t, J=6.8Hz), 3.98 (3H, s), 7.50 (1H, dd, J=1.5 and 8.5Hz), 7.79 (1H, d, J=8.6Hz), 7.85 (1H, d, J=8.5Hz), 7.92 (1H, s), 8.04 (1H, dd, J=1.7 and 8.6Hz), 8.55 (1H, s)

APCI-MASS : m/z = 267 (M⁺+1)

10 Preparation 291

To a solution of 4-octylaniline (5 ml) in a mixture of pyridine (12.5 ml) and chloroform (40 ml) was added phenyl chloroformate (2.95 ml) and stirred for 1.5 hours at ambient temperature. The reaction mixture was added to a mixture of water and ethyl acetate. The organic layer was taken and dried over magnesium sulfate. The magnesium sulfate was filtered off, and the filtrate was evaporated under reduced pressure to give 4-Octyl-N-phenoxy carbonylaniline (4.51 g)

IR (KBr) : 3318.9, 1714.4, 1234.2 cm⁻¹

20 NMR (CDCl₃, δ) : 0.88 (3H, t, J=6.2Hz), 1.2-1.4 (10H, m), 1.5-1.7 (2H, m), 2.57 (2H, t, J=7.3Hz), 6.88 (1H, s), 7.1-7.5 (9H, m)

25 The following compounds (Preparations 292 to 299) were obtained according to a similar manner to that of Preparation 291.

Preparation 292

30 4-(4-tert-Butoxycarbonylpiperazin-1-yl)-N-phenoxy carbonylaniline

IR (KBr) : 3309.2, 1743.3, 1658.5, 1197.6 cm⁻¹

NMR (CDCl₃, δ) : 1.48 (9H, s), 3.08 (4H, t, J=5.3Hz), 3.58 (4H, t, J=5.3Hz), 6.87 (1H, s), 6.91 (2H, d, J=9Hz), 7.1-7.5 (7H, m)

35 APCI-MASS : m/z = 398 (M+H⁺)

Preparation 293

1-(4-Cyclohexylbenzoyl)-2-(4-methoxycarbonylbenzoyl)-hydrazine

IR (KBr) : 3236, 2925, 2852, 1726, 1679, 1637, 1278,
5 1110 cm^{-1}

NMR (DMSO-d_6 , δ) : 1.1-1.5 (5H, m), 1.6-2.0 (5H, m),
2.60 (1H, m), 3.90 (3H, s), 7.37 (2H, d, $J=8.0\text{Hz}$),
7.85 (2H, d, $J=8.0\text{Hz}$), 8.0-8.2 (4H, m), 10.48 (1H,
s), 10.68 (1H, s)

10 APCI-MASS : $m/z = 381 \text{ (M+H)}^+$

Preparation 294

1-[4-(Piperidin-1-yl)benzoyl]-2-(4-methoxycarbonylbenzoyl)hydrazine

15 IR (KBr) : 3500, 3286, 2941, 2854, 1712, 1689, 1650,
1606, 1286, 1242 cm^{-1}

NMR (DMSO-d_6 , δ) : 1.59 (6H, s), 3.33 (4H, s), 3.90
(3H, s), 6.97 (2H, d, $J=8.8\text{Hz}$), 7.79 (2H, d,
 $J=8.8\text{Hz}$), 8.02 (2H, d, $J=8.4\text{Hz}$), 8.09 (2H, d,
20 $J=8.4\text{Hz}$), 10.23 (1H, s), 10.57 (1H, s)

APCI-MASS : $m/z = 382 \text{ (M+H)}^+$

Preparation 295

25 1-[4-(4-n-Propyloxyphenyl)benzoyl]-2-(4-methoxycarbonylbenzoyl)hydrazine

IR (KBr) : 3230, 1724, 1679, 1654, 1280, 1108 cm^{-1}

NMR (DMSO-d_6 , δ) : 1.00 3H, d, $J=7.5\text{Hz}$), 1.76 (2H, tq,
 $J=6.5$ and 7.5Hz), 3.91 (3H, s), 7.05 (2H, d,
 $J=8.7\text{Hz}$), 7.71 (2H, d, $J=8.7\text{Hz}$), 7.79 (2H, d,
30 $J=8.5\text{Hz}$), 8.00 (2H, d, $J=8.5\text{Hz}$), 8.05 (2H, d,
 $J=8.6\text{Hz}$), 8.11 (2H, d, $J=8.6\text{Hz}$), 10.60 (1H, s),
10.72 (1H, s)

APCI-MASS : $m/z = 433 \text{ (M+H)}^+$

35 Preparation 296

1-(4-Methoxycarbonylbenzoyl)-2-decanoylhydrazine

IR (KBr) : 3220, 2919, 2850, 1724, 1643, 1600, 1567,
1479, 1284 cm^{-1}

NMR (DMSO- d_6 , δ) : 0.86 (3H, t, $J=6.8\text{Hz}$), 1.2-1.7
(14H, m), 2.18 (2H, t, $J=7.4\text{Hz}$), 3.89 (3H, s), 7.97
(2H, d, $J=8.5\text{Hz}$), 8.06 (2H, d, $J=8.5\text{Hz}$), 9.15 (1H,
s), 10.49 (1H, s)

APCI-MASS : $m/z = 349$ ($\text{M}+\text{H}^+$)

10 Preparation 297

1-(4-Methoxycarbonylbenzoyl)-2-(trans-4-n-
pentylcyclohexylcarbonyl)hydrazine

IR (KBr) : 3201, 2923, 2852, 1727, 1600, 1567, 1479,
1282 cm^{-1}

15 NMR (DMSO- d_6 , δ) : 0.86 (3H, t, $J=6.9\text{Hz}$), 0.9-1.0 (2H,
m), 1.1-1.5 (11H, m), 1.7-1.9 (4H, m), 2.20 (1H,
m), 3.88 (3H, s), 7.97 (2H, d, $J=8.6\text{Hz}$), 8.06 (2H,
d, $J=8.6\text{Hz}$), 9.85 (1H, s), 10.46 (1H, s)

APCI-MASS : $m/z = 375$ ($\text{M}+\text{H}^+$)

20

Preparation 298

1-[4-(8-Methoxyoctyloxy)benzoyl]-2-(4-
methoxycarbonylbenzoyl)hydrazine

IR (KBr) : 3213, 2935, 2856, 1718, 1600, 1567, 1465,
1282 cm^{-1}

25 NMR (DMSO- d_6 , δ) : 1.2-1.8 (12H, m), 3.21 (3H, s),
3.29 (2H, t, $J=6.4\text{Hz}$), 3.90 (3H, s), 4.04 (2H, t,
 $J=6.5\text{Hz}$), 7.04 (2H, d, $J=8.8\text{Hz}$), 7.90 (2H, d,
 $J=8.8\text{Hz}$), 8.04 (2H, d, $J=8.7\text{Hz}$), 8.10 (2H, d,
30 $J=8.7\text{Hz}$), 10.41 (1H, s), 10.64 (1H, s)

APCI-MASS : $m/z = 457$ ($\text{M}+\text{H}^+$)

Preparation 299

35 1-(4-Octyloxybenzoyl)-2-(4-methoxycarbonylbenzoyl)-
hydrazine

0360972 0344 0360980

IR (KBr) : 3224, 2923, 2854, 1724, 1681, 1643, 1502,
1434, 1282, 1253, 1106 cm^{-1}

NMR (DMSO- d_6 , δ) : 0.86 (3H, t, $J=6.8\text{Hz}$), 1.2-1.5
(10H, m), 1.6-1.8 (2H, m), 3.89 (3H, s), 4.04 (2H,
5 t, $J=6.3\text{Hz}$), 7.04 (2H, d, $J=8.7\text{Hz}$), 7.90 (2H, d,
 $J=8.7\text{Hz}$), 8.03 (2H, d, $J=8.6\text{Hz}$), 8.10 (2H, d,
 $J=8.6\text{Hz}$), 10.42 (1H, s), 10.64 (1H, s)

APCI-MASS : $m/z = 427$ ($\text{M}+\text{H}^+$)

10 Preparation 300

A solution of Methyl 4-n-hexyloxybenzoate (2.00 g) and
hydrazine hydrate (4.24 g) in ethanol (10 ml) was refluxed
for 6 hours. After cooling, the reaction mixture was poured
into water. The precipitate was collected by filtration,
15 washed with water and dried over P_2O_5 under reduced pressure
to give N-(4-n-hexyloxybenzoyl)hydrazine (1.96 g).

IR (KBr) : 3311, 2954, 2869, 1623, 1253 cm^{-1}

NMR (DMSO- d_6 , δ) : 0.87 (3H, t, $J=6.8\text{Hz}$), 1.2-1.5 (6H,
m), 1.6-1.8 (2H, m), 4.00 (2H, t, $J=6.5\text{Hz}$), 4.40
20 (2H, s), 6.95 (2H, d, $J=8.6\text{Hz}$), 7.77 (2H, d,
 $J=8.6\text{Hz}$), 9.59 (1H, s)

APCI-MASS : $m/z = 237$ ($\text{M}+\text{H}^+$)

The following compounds (Preparations 301 to 308) were
25 obtained according to a similar manner to that of Preparation
300.

Preparation 301

N-(4-Octylphenyl)-N'-aminourea

30 IR (KBr) : 3309.2, 1683.6, 1554.3 cm^{-1}

NMR (DMSO- d_6 , δ) : 0.85 (3H, t, $J=6.7\text{Hz}$), 1.1-1.4
(10H, m), 1.4-1.6 (2H, m), 2.48 (2H, t, $J=8.9\text{Hz}$),
4.32 (2H, s), 7.03 (2H, d, $J=8.4\text{Hz}$), 7.32 (1H, s),
7.38 (2H, d, $J=8.4\text{Hz}$), 8.50 (1H, s)

Preparation 302

N-[4-(4-tert-Butoxycarbonylpiperazin-1-yl)phenyl]-N'-aminourea

IR (KBr) : 3237.9, 1695.1, 1670.1, 1540.8, 1230.4 cm^{-1}

5 NMR (DMSO- d_6 , δ) : 1.42 (9H, s), 2.97 (4H, t, J=4.9Hz), 3.44 (4H, t, J=4.9Hz), 4.30 (2H, s), 6.85 (2H, d, J=9.0Hz), 7.26 (1H, s), 7.36 (2H, d, J=9.0Hz), 8.41 (1H, s)

10 Preparation 303

4-Cyclohexylbenzoylhydrazine

IR (KBr) : 3318, 2925, 2852, 1625, 1606, 1527, 1326 cm^{-1}

15 NMR (DMSO- d_6 , δ) : 1.1-1.5 (5H, m), 1.6-2.0 (5H, m), 2.4-2.6 (1H, m), 4.44 (2H, s), 7.27 (2H, d, J=8.2Hz), 7.73 (2H, d, J=8.2Hz), 9.66 (1H, s)

APCI-MASS : $m/z = 219$ (M+H)⁺

Preparation 304

20 4-(Piperidin-1-yl)benzoylhydrazine

IR (KBr) : 3263, 2852, 1612, 1504, 1245, 1124 cm^{-1}

NMR (DMSO- d_6 , δ) : 1.57 (6H, s), 3.25 (4H, s), 4.35 (2H, s), 6.90 (2H, d, J=9.0Hz), 7.68 (2H, d, J=9.0Hz), 9.44 (1H, s)

25 APCI-MASS : $m/z = 220$ (M+H)⁺

Preparation 305

4-(4-n-Propyloxyphenyl)benzoylhydrazine

IR (KBr) : 3350, 3276, 1610, 1494, 1288, 978 cm^{-1}

30 NMR (DMSO- d_3 , δ) : 0.99 (3H, t, J=7.5Hz), 1.75 (2H, tq, J=6.5 and 7.5Hz), 3.98 (2H, t, J=6.5Hz), 4.50 (2H, s), 7.03 (2H, d, J=8.8Hz), 7.65 (2H, d, J=8.8Hz), 7.69 (2H, d, J=8.4Hz), 7.88 (2H, d, J=8.4Hz), 9.79 (1H, s)

35 APCI-MASS : $m/z = 271$ (M+H)⁺

Preparation 306

4-Methoxycarbonylbenzoylhydrazine

IR (KBr) : 3322, 3250, 3018, 1727, 1658, 1621, 1565,
1432, 1280, 1110 cm^{-1}

5 NMR (DMSO- d_6 , δ) : 3.87 (3H, s), 4.58 (2H, s), 7.93
(2H, dd, $J=8.6$ and 3.1Hz), 7.02 (2H, dd, $J=8.6$ and
3.1Hz), 9.97 (1H, s)

APCI-MASS : $m/z = 195$ ($\text{M}+\text{H}^+$)

10 Preparation 307

Trans-4-n-pentylcyclohexylcarbonylhydrazine

IR (KBr) : 3303, 3199, 2954, 2925, 2850, 1639, 1619,
1533, 1457 cm^{-1}

15 NMR (DMSO- d_6 , δ) : 0.8-1.0 (6H, m), 1.1-1.5 (10H, m),
1.6-2.2 (5H, m), 4.10 (2H, s), 8.85 (1H, s)

APCI-MASS : $m/z = 213$ ($\text{M}+\text{H}^+$)

Preparation 308

4-(8-Methoxyoctyloxy)benzoylhydrazine

20 IR (KBr) : 3309, 2937, 2852, 1606, 1494, 1253 cm^{-1}

NMR (DMSO- d_6 , δ) : 1.2-1.8 (12H, m), 3.20 (3H, s),
3.25 (2H, t, $J=6.5\text{Hz}$), 3.99 (2H, t, $J=6.5\text{Hz}$), 4.39
(2H, s), 6.95 (2H, d, $J=8.8\text{Hz}$), 7.77 (2H, d,
 $J=8.8\text{Hz}$), 9.58 (1H, s)

25 APCI-MASS : $m/z = 295$ ($\text{M}+\text{H}^+$)

Preparation 309

To a stirred solution of 4-bromo-4'-n-heptylbiphenyl
(2.71 g) in tetrahydrofuran (100 ml) was added dropwise a
30 solution of n-butyllithium in a mixture of diethyl ether and
n-hexane (1.6M, 5.1 ml) at -78°C . After stirring at -78°C
for 30 minutes, the resultant mixture was added to a solution
of diethyl oxalate (3.4 ml) in tetrahydrofuran (50 ml) at
 -78°C . The resultant mixture was allowed to warm to 0°C for
35 about 1 hour, and to the mixture was added acetic acid (0.5

ml). Evaporation gave a residue which was taken up into a mixture of water and ethyl acetate. The organic layer was separated, washed with brine, dried over magnesium sulfate. Evaporation gave a residue which was chromatographed on silica gel (200 ml) eluting with a mixture of n-hexane and ethyl acetate (10:0-95:5, V/V) to give 1-Ethyl-2-(4-n heptylphenyl)ethanedione (2.23 g).

NMR (CDCl₃, δ) : 0.88 (3H, t, J=6.6Hz), 1.10-1.50 (8H, m), 1.44 (3H, t, J=7.1Hz), 1.50-1.80 (2H, m), 2.66 (2H, t, J=7.7Hz), 4.47 (2H, q, J=7.1Hz), 7.20-7.40 (2H, m), 7.50-7.64 (2H, m), 7.64-7.85 (2H, m), 8.00-8.20 (2H, m)

APCI-MASS : m/z = 353 (M⁺+1)

Preparation 310

To a suspension of sodium hydride (60% in oil, 0.37 g) in tetrahydrofuran (40 ml) was added by portions 4-acetyl-4'-n-heptylbiphenyl (2.50 g) at ambient temperature. After stirring at ambient temperature for 1 hour, to the solution was added triethyl phosphonoacetate (1.9 ml) and the mixture was heated to reflux for 5 hours. After cooling to ambient temperature, to the mixture was added acetic acid (0.53 ml) and evaporated. The residue was taken up into a mixture of water and ethyl acetate. The separated organic layer was washed with brine, dried over magnesium sulfate and evaporated. The residue was chromatographed on silica gel (200 ml) eluting with mixture of n-hexane and diisopropyl ether (99:1-20:1, V/V) to give Ethyl (E)-3-[4-(4-heptylphenyl)phenyl]-2-butenolate (2.19 g).

NMR (CDCl₃, δ) : 0.88 (3H, t, J=6.6Hz), 1.13-1.48 (8H, m), 1.48-1.78 (2H, m), 2.61 (3H, s), 2.65 (2H, t, J=7.4Hz), 4.22 (2H, q, J=7.1Hz), 6.20 (1H, t, J=2.7Hz), 7.23-7.28 (2H, m), 7.50-7.63 (6H, m)

APCI-MASS : m/z = 365 (M⁺+1)

Preparation 311

To a solution of 4-bromo-4'-n-heptylbiphenyl (5.1 g) in tetrahydrofuran (60 ml) was added a solution of n-butyllithium in a mixture of n-hexane and diethyl ether (1.6M, 9.7 ml) at -60°C. After stirring at -60°C for 30 minutes, to the mixture was added N,N-dimethylacetamide (4.3 ml) and the reaction mixture was allowed to warm to 0°C. The reaction mixture was taken up into a mixture of cold water and ethyl acetate, and the pH was adjusted to around 1 with 1N hydrochloric acid. The organic layer was separated, washed with brine, dried over magnesium sulfate and evaporated. The residue was chromatographed on silica gel (150 ml) eluting with a mixture of n-hexane and ethyl acetate (20:1, V/V) to give 4-Acetyl-4'-n-heptylbiphenyl (1.60 g).

NMR (CDCl₃, δ) : 0.89 (3H, t, J=6.6Hz), 1.05-1.48 (8H, m), 1.48-1.75 (2H, m), 2.65 (2H, t, J=7.6Hz), 2.63 (3H, s), 7.20-7.31 (2H, m), 7.52-7.58 (2H, m), 7.65-7.70 (2H, m), 7.97-8.05 (2H, m)

APCI-MASS : m/z = 295 (M+1)

Preparation 312

To a solution of Methyl 4-[4-(8-hydroxyoctyloxy)phenyl]-benzoate (500 mg) and dihydropyran (141 mg) in dichloromethane (15 ml) was added p-toluenesulfonic acid (5 ml). The mixture was stirred at ambient temperature for 10 minutes and diluted with dichloromethane and washed with water and brine. The separated organic layer was dried over magnesium sulfate and evaporated under reduced pressure to give Methyl 4-[4-(8-tetrahydropyran-2-yl-oxyoctyloxy)phenyl]-benzoate (616 mg).

IR (KBr) : 2935, 2856, 1722, 1602, 1438, 1290, 1199 cm⁻¹

NMR (CDCl₃, δ) : 1.3-2.0 (18H, m), 3.3-3.9 (4H, m), 3.93 (3H, s), 4.00 (2H, t, J=6.5Hz), 4.5-4.6 (1H, m), 6.98 (2H, d, J=8.7Hz), 7.56 (2H, d, J=8.7Hz),

7.62 (2H, d, J=8.3Hz), 8.07 (2H, d, J=8.3Hz)

Preparation 313

To a solution of titanium(IV) chloride (11.6 g) in
5 dichloromethane (100 ml) was added 4-n-Pentyloxyacetophenone
(10.3 g) and Methyl 4-formylbenzoate (8.21 g) in
dichloromethane (50 ml) dropwise at 0°C. To the mixture was
added triethylamine (11.15 ml) in dichloromethane (30 ml).
The mixture was stirred at 0°C for 30 minutes and diluted
10 with n-hexane. The organic layer was washed with water (four
times), brine and dried over magnesium sulfate. The solvents
were removed under reduced pressure and the residue was
trituated with iso-propyl ether. The solid was collected by
filtration and dried to give 1-(4-Methoxycarbonylphenyl)-3-
15 (4-n-pentyloxyphenyl)-1-propen-3-one (4.02 g).

IR (KBr) : 2950, 2910, 2863, 1718, 1654, 1606, 1274,
1176 cm⁻¹

NMR (CDCl₃, δ) : 0.94 (3H, t, J=6.9Hz), 1.3-1.6 (4H,
m), 1.8-2.0 (2H, m), 3.93 (3H, s), 4.04 (2H, t,
20 J=6.5Hz), 6.97 (2H, d, J=8.8Hz), 7.60 (1H, d,
J=15.7Hz), 7.68 (2H, d, J=8.4Hz), 7.80 (1H, d,
J=15.7Hz), 8.0-8.2 (4H, m)

APCI-MASS : m/z = 353 (M+H⁺)

25 Preparation 314

To a solution of titanium(IV) chloride (13.88 g) in
dichloromethane (100 ml) was added Ethyl 4-acetylbenzoate
(11.53 g) and 4-n-pentyloxybenzaldehyde (12.69 g) in
dichloromethane (50 ml) was added dropwise at 0°C. To the
30 mixture was added triethylamine (12.44 ml) in dichloromethane
(30 ml). The mixture was stirred at 0°C for 30 minutes and
diluted with ethyl acetate. The organic layer was washed
with water (four times) and brine and dried over magnesium
sulfate. The solvents were removed under reduced pressure
35 and the residue was trituated with n-hexane. The solid was

collected by filtration and dried to give 1-(4-n-Pentyloxyphenyl)-3-(4-ethoxycarbonylphenyl)-1-propen-3-one (13.45 g).

IR (KBr) : 2956, 2929, 2861, 1718, 1656, 1594, 1510, 1272 cm^{-1}

NMR (CDCl_3 , δ) : 0.94 (3H, t, $J=7.1\text{Hz}$), 1.3-1.9 (9H, m), 4.01 (2H, t, $J=6.5\text{Hz}$), 4.42 (2H, q, $J=7.1\text{Hz}$), 6.93 (1H, d, $J=8.7\text{Hz}$), 7.37 (1H, d, $J=15.6\text{Hz}$), 7.60 (2H, d, $J=8.7\text{Hz}$), 7.81 (1H, d, $J=15.6\text{Hz}$), 8.03 (2H, d, $J=8.5\text{Hz}$), 8.16 (2H, d, $J=8.5\text{Hz}$)

APCI-MASS : $m/z = 367$ ($M+H^+$)

The following compound was obtained according to a similar manner to that of Preparation 314.

Preparation 315

Ethyl 4-oxo-1-(4-n-hexyloxyphenyl)piperidine-3-carboxylate

IR (Neat) : 1664.3, 1511.9, 1243.9, 1216.9 cm^{-1}

NMR (CDCl_3 , δ) : 0.90 (3H, t, $J=6.5\text{Hz}$), 1.2-1.5 (6H, m), 1.32 (3H, t, $J=7.1\text{Hz}$), 1.65-1.85 (2H, m), 2.51 (2H, t, $J=5.8\text{Hz}$), 3.31 (2H, t, $J=5.8\text{Hz}$), 3.76 (2H, s), 3.91 (2H, t, $J=6.5\text{Hz}$), 4.26 (2H, q, $J=7.1\text{Hz}$), 6.84 (2H, d, $J=9.2\text{Hz}$), 6.94 (2H, d, $J=9.2\text{Hz}$), 12.06 (1H, s)

APCI-MASS : $m/z = 348$ (M^++H)

Preparation 316

To a solution of 4-n-Hexyloxybenzoylhydrazine (1.96 g) and pyridine (0.74 ml) in tetrahydrofuran (20 ml) was added a solution of terephthalic acid monomethyl ester chloride (1.56 g) in tetrahydrofuran (15 ml) dropwise at 0°C . The reaction mixture was stirred at room temperature for 2 hours, and poured into water. The precipitate was collected by filtration and washed with acetonitrile. The residue was

dried under reduced pressure to give 1-(4-n-Hexyloxybenzoyl)-
2-(4-methoxycarbonylbenzoyl)hydrazine (2.99 g).

IR (KBr) : 3230, 3023, 2954, 2858, 1724, 1681, 1643,
1280, 1251, 1105 cm^{-1}

NMR (DMSO-d_6 , δ) : 0.88 (3H, t, $J=6.6\text{Hz}$), 1.2-1.5 (6H,
m), 1.6-1.8 (2H, m), 3.90 (3H, s), 4.04 (2H, t,
 $J=6.4\text{Hz}$), 7.04 (2H, d, $J=8.7\text{Hz}$), 7.90 (2H, d,
 $J=8.7\text{Hz}$), 8.03 (2H, d, $J=8.4\text{Hz}$), 8.10 (2H, d,
 $J=8.4\text{Hz}$), 10.42 (1H, s), 10.65 (1H, s)

APCI-MASS : $m/z = 399 \text{ (M+H)}^+$

Preparation 317

A mixture of 1-(4-n-Hexyloxyphenyl)-4-piperidone (0.823
g), 1-(4-Ethoxycarbonylphenyl)piperazine (0.7 g), and
titanium(IV) isopropoxide (1.11 ml) was stirred at room
temperature. After 1 hour, the IR spectrum of the mixture
showed no ketone band, and the viscous solution was diluted
with absolute ethanol (3 ml). Sodium cyanoborohydride (0.121
g) was added, and the solution was stirred for 3 hours.
Water (3 ml) was added with stirring, and the resulting in
organic precipitate was filtered and washed with ethanol.
The filtrate was extracted with ethyl acetate. The organic
layer was taken and dried over magnesium sulfate. The
magnesium sulfate was filtered off, and filtrate was
evaporated under reduced pressure to give Ethyl 4-[4-[1-(4-n-
hexyloxyphenyl)piperidin-4-yl]piperazin-1-yl]benzoate (331
mg).

IR (KBr) : 1708.6, 1606.4, 1511.9, 1284.4, 1236.1 cm^{-1}

NMR (CDCl_3 , δ) : 0.90 (3H, t, $J=6.5\text{Hz}$), 1.2-1.55 (6H,
m), 1.37 (3H, t, $J=7.1\text{Hz}$), 1.6-1.85 (4H, m), 1.95
(2H, d, $J=12\text{Hz}$), 2.41 (1H, m), 2.62 (2H, d,
 $J=11\text{Hz}$), 2.75 (4H, t, $J=5.0\text{Hz}$), 3.35 (4H, t,
 $J=5.0\text{Hz}$), 3.58 (2H, d, $J=11\text{Hz}$), 3.90 (2H, t,
 $J=6.5\text{Hz}$), 4.32 (2H, q, $J=7.1\text{Hz}$), 6.7-7.0 (6H, m),
7.92 (2H, d, $J=9.0\text{Hz}$)

APCI-MASS : $m/z = 494$ ($M^+ + H$)

The following compound was obtained according to a similar manner to that of Preparation 317.

Preparation 318

1-tert-Butoxycarbonyl-4-(4-phenylcyclohexyl)piperazine

IR (KBr) : 1697.1, 1245.8, 1170.6, 1124.3, 700 cm^{-1}

NMR (CDCl_3 , δ) : 1.2-1.65 (17H, m), 1.9-2.1 (4H, m),
2.3-2.6 (2H, m), 2.55 (4H, t, $J=5.0\text{Hz}$), 3.44 (4H,
t, $J=5.0\text{Hz}$), 7.1-7.4 (5H, m)

APCI-MASS : $m/z = 345$ ($M^+ + H$)

Preparation 319

To a suspension of 1-(N,N-dimethylamino)-2-(4-ethoxycarbonylbenzoyl)ethylene (0.742 g) and 4-n-hexyloxybenzamidinium hydrochloride (0.847 g) in methanol (10 ml) was added 28% sodium methoxide in methanol (0.64 ml). The suspension was refluxed for 6 hours, and partitioned with ethyl acetate and water. The organic layer was washed with water and brine, dried over magnesium sulfate and evaporated under reduced pressure. The residue was triturated with acetonitrile, collected by filtration and dried under reduced pressure to give Methyl 4-[2-(4-n-hexyloxyphenyl)pyrimidin-6-yl]benzoate (0.61 g).

IR (KBr) : 2931, 2861, 1722, 1606, 1558, 1251 cm^{-1}

NMR (CDCl_3 , δ) : 0.95 (3H, t, $J=6.7\text{Hz}$), 1.2-1.6 (6H, m), 1.8-2.0 (2H, m), 3.97 (3H, s), 4.05 (2H, t, $J=6.5\text{Hz}$), 7.02 (2H, d, $J=8.8\text{Hz}$), 7.56 (1H, d, $J=5.2\text{Hz}$), 8.18 (2H, d, $J=8.6\text{Hz}$), 8.28 (2H, d, $J=8.6\text{Hz}$), 8.52 (2H, d, $J=8.8\text{Hz}$), 8.83 (1H, d, $J=5.2\text{Hz}$)

APCI-MASS : $m/z = 391$ ($M + H^+$)

Preparation 320

A solution of 1-(4-Methoxycarbonylphenyl)-3-(4-n-pentyloxyphenyl)-1-propen-3-one (4.0 g) and hydroxyamine hydrochloride (3.93 g) in ethanol (40 ml) was refluxed for 4 hours. The mixture was diluted with ethyl acetate, and the organic layer was washed with water (x 2), brine and dried over magnesium sulfate. The solvents were removed under reduced pressure to give crude oxime. To a solution of crude oxime in 1,2-dichloroethane (20 ml) was added activated-manganese(IV) oxide (10.0 g). The reaction mixture was refluxed for 2 hours and filtered. The residue was washed with dichloromethane. The solvents were removed under reduced pressure and the residue was triturated with acetonitrile. The solid was collected by filtration and dried to give Methyl 4-[3-(4-n-pentyloxyphenyl)isoxazol-5-yl]benzoate (0.98 g).

IR (KBr) : 2940, 2871, 1720, 1612, 1278, 1249, 1178, 1108 cm^{-1}

NMR (DMSO-d_6 , δ) : 0.94 (3H, t, $J=7.2\text{Hz}$), 1.2-1.6 (4H, m), 1.7-1.9 (2H, m), 3.95 (3H, s), 4.01 (2H, t, $J=6.5\text{Hz}$), 6.87 (1H, s), 6.98 (2H, d, $J=8.9\text{Hz}$), 7.79 (2H, d, $J=8.9\text{Hz}$), 7.89 (2H, d, $J=8.6\text{Hz}$), 8.15 (2H, d, $J=8.6\text{Hz}$)

APCI-MASS : $m/z = 366$ ($\text{M}+\text{H}^+$)

Preparation 321

To a solution of 4-Methoxycarbonylphenylhydroxyimino-methyl chloride (16.98 g) and 4-n-pentyloxyphenylacetylene (18.96 g) in tetrahydrofuran (170 ml) was added triethylamine (14.4 ml) in tetrahydrofuran (140 ml) over a period of 2 hours at 40°C and the mixture was stirred at 40°C for 30 minutes. The mixture was diluted with dichloromethane and washed with water and brine. The separated organic layer was dried over magnesium sulfate and evaporated under reduced pressure. The residue was triturated with acetonitrile. The precipitate was collected by filtration and dried to give

Methyl 4-[5-(4-n-pentyloxyphenyl)isoxazol-3-yl]benzoate
(24.56 g).

IR (KBr) : 2942, 2873, 1716, 1616, 1508, 1280,
1108 cm^{-1}

5 NMR (CDCl_3 , δ) : 0.95 (3H, t, $J=6.9\text{Hz}$), 1.3-1.6 (4H,
m), 1.8-2.0 (2H, m), 3.95 (3H, s), 4.02 (2H, t,
 $J=6.5\text{Hz}$), 6.74 (1H, s), 6.99 (2H, d, $J=8.8\text{Hz}$), 7.76
(2H, d, $J=8.8\text{Hz}$), 7.93 (2H, d, $J=8.5\text{Hz}$), 8.14 (2H,
d, $J=8.5\text{Hz}$)

10 APCI-MASS : m/z = 366 ($\text{M}+\text{H}^+$)

Preparation 322

467250-22/60899
To a solution of N-Hydroxy-4-octyloxybenzamidine (1.89
g) in pyridine (10 ml) was added terephthalic acid monomethyl
15 ester chloride (1.67 g) in tetrahydrofuran (15 ml) dropwise
at 0°C. The mixture was stirred at room temperature for 15
minutes, and poured into water. The precipitate was
collected by filtration, dried and dissolved in pyridine (10
ml). The solution was refluxed for 1 hour. The reaction
20 mixture was diluted with ethyl acetate and washed with 1N
HCl, water and brine. The separated organic layer was dried
over magnesium sulfate and the solvents were removed under
reduced pressure. The residue was triturated with
acetonitrile and collected by filtration. The solid was
25 dried to give Methyl 4-[3-(4-n-hexyloxyphenyl)-1,2,4-
oxadiazol-5-yl]benzoate (2.27 g).

IR (KBr) : 2950, 2925, 2863, 1720, 1280, 1255 cm^{-1}

NMR (CDCl_3 , δ) : 0.92 (3H, t, $J=6.6\text{Hz}$), 1.2-1.9 (8H,
m), 3.97 (3H, s), 4.03 (2H, d, $J=6.5\text{Hz}$), 7.00 (2H,
30 d, $J=8.9\text{Hz}$), 8.09 (2H, d, $J=8.9\text{Hz}$), 8.20 (2H, d,
 $J=6.6\text{Hz}$), 8.28 (2H, d, $J=6.6\text{Hz}$)

APCI-MASS : m/z = 381 ($\text{M}+\text{H}^+$)

Preparation 323

35 A suspension of 1-(4-n-Hexyloxybenzoyl)-2-(4-

methoxycarbonylbenzoyl)hydrazine (1.00 g) in phosphorus
oxychloride (5 ml) was refluxed for 1 hour. After cooling,
the solution was concentrated under reduced pressure. The
residue was poured into ice-water and extracted with
5 dichloromethane. The organic layer was washed with water,
brine and dried over magnesium sulfate. The solvents were
removed under reduced pressure. The residue was triturated
with acetonitrile, collected by filtration and dried under
reduced pressure to give Methyl 4-[5-(4-n-hexyloxyphenyl)-
10 1,3,4-oxadiazole-2-yl]benzoate (761 mg).

IR (KBr) : 2954, 2854, 1724, 1612, 1494, 1280,
1249 cm^{-1}

NMR (CDCl_3 , δ) : 0.91 (3H, t, $J=6.6\text{Hz}$), 1.3-1.6 (6H,
m), 1.7-1.9 (2H, m), 3.96 (3H, s), 4.04 (2H, t,
15 $J=6.5\text{Hz}$), 7.02 (2H, d, $J=8.6\text{Hz}$), 8.07 (2H, d,
 $J=8.6\text{Hz}$), 8.19 (4H, m)

APCI-MASS : $m/z = 381$ ($M+H$)⁺

The following compounds (Preparations 324 to 327) were
20 obtained according to a similar manner to that of Preparation
323.

Preparation 324

Methyl 4-[5-[4-(4-n-propyloxyphenyl)phenyl]-1,3,4-
25 oxadiazol-2-yl]benzoate

IR (KBr) : 1720, 1614, 1496, 1280, 1103 cm^{-1}

NMR (CDCl_3 , δ) : 1.07 (3H, d, $J=7.5\text{Hz}$), 1.84 (2H, tq,
 $J=6.5$ and 7.5Hz), 3.98 (3H, s), 3.99 (2H, t,
 $J=6.5\text{Hz}$), 7.01 (2H, d, $J=8.8\text{Hz}$), 7.60 (2H, d,
30 $J=8.8\text{Hz}$), 7.73 (2H, d, $J=8.5\text{Hz}$), 8.19 (2H, d,
 $J=8.5\text{Hz}$), 8.22 (4H, s)

APCI-MASS : $m/z = 415$ ($M+H$)⁺

Preparation 325

35 Methyl 4-[5-(n-nonyl)-1,3,4-oxadiazol-2-yl]benzoate

IR (KBr) : 2915, 2848, 1724, 1569, 1436, 1413,
1278 cm^{-1}

NMR (CDCl_3 , δ) : 0.88 (3H, t, $J=6.4\text{Hz}$), 1.2-1.6 (12H, m), 1.8-2.0 (2H, m), 2.94 (2H, t, $J=7.6\text{Hz}$), 3.96 (3H, s), 8.11 (2H, d, $J=8.8\text{Hz}$), 8.17 (2H, d, $J=8.8\text{Hz}$)

APCI-MASS : $m/z = 331$ ($\text{M}+\text{H}^+$)

Preparation 326

Methyl 4-[5-[4-(8-methoxyoctyloxy)phenyl]-1,3,4-oxadiazol-2-yl]benzoate

IR (KBr) : 2925, 2858, 1722, 1614, 1280, 1259 cm^{-1}

NMR (CDCl_3 , δ) : 1.3-1.9 (12H, m), 3.36 (3H, s), 3.37 (2H, t, $J=6.4\text{Hz}$), 3.97 (3H, s), 4.04 (2H, t, $J=6.5\text{Hz}$), 7.02 (2H, d, $J=8.9\text{Hz}$), 8.07 (2H, d, $J=8.9\text{Hz}$), 8.20 (4H, s)

APCI-MASS : $m/z = 439$ ($\text{M}+\text{H}^+$)

Preparation 327

Methyl 4-[5-(4-n-octyloxyphenyl)-1,3,4-oxadiazol-2-yl]benzoate

IR (KBr) : 2923, 2856, 1722, 1614, 1496, 1282,
1103 cm^{-1}

NMR (CDCl_3 , δ) : 0.89 (3H, t, $J=6.8\text{Hz}$), 1.2-1.6 (10H, m), 1.7-1.9 (2H, m), 3.97 (3H, s), 4.04 (2H, t, $J=6.5\text{Hz}$), 7.03 (2H, d, $J=8.7\text{Hz}$), 8.07 (2H, d, $J=8.7\text{Hz}$), 8.19 (4H, m)

APCI-MASS : $m/z = 409$ ($\text{M}+\text{H}^+$)

Preparation 328

A suspension of 1-(4-Hexyloxybenzoyl)-2-(4-methoxycarbonylbenzoyl)hydrazine (1.0 g) and di-phosphorus pentasulfide (1.28 g) in tetrahydrofuran (15 ml) was stirred at room temperature for 3 hours. The mixture was diluted with water (30 ml), stirred for 30 minutes and extracted with

dichloromethane. The organic layer was washed with brine, dried over magnesium sulfate and evaporated under reduced pressure. The residue was triturated with acetonitrile. The solid was collected by filtration and dried under reduced pressure to give Methyl 4-[5-(4-n-hexyloxyphenyl)-1,3,4-thiadiazol-2-yl]benzoate (816 mg).

IR (KBr) : 2925, 2871, 1722, 1608, 1436, 1276,
1106 cm^{-1}

NMR (CDCl_3 , δ) : 0.92 (3H, t, $J=6.6\text{Hz}$), 1.3-2.0 (8H, m), 3.96 (3H, s), 4.03 (2H, t, $J=6.5\text{Hz}$), 6.99 (2H, d, $J=8.6\text{Hz}$), 7.95 (2H, d, $J=8.4\text{Hz}$), 8.16 (2H, d, $J=8.4\text{Hz}$)

APCI-MASS : $m/z = 397$ ($M+H$)⁺

The following compounds (Preparations 329 to 334) were obtained according to a similar manner to that of Preparation 328.

Preparation 329

Methyl 4-[5-[4-(8-methoxyoctyloxy)phenyl]-1,3,4-thiadiazol-2-yl]benzoate

IR (KBr) : 3210, 2935, 2856, 1718, 1600, 1465, 1280,
1110 cm^{-1}

NMR (CDCl_3 , δ) : 1.3-1.6 (10H, m), 1.7-1.9 (2H, m), 3.33 (3H, s), 3.37 (2H, d, $J=6.4\text{Hz}$), 3.96 (3H, s), 4.03 (2H, t, $J=6.5\text{Hz}$), 6.99 (2H, d, $J=8.9\text{Hz}$), 7.94 (2H, d, $J=8.9\text{Hz}$), 8.07 (2H, d, $J=8.6\text{Hz}$), 8.16 (2H, d, $J=8.6\text{Hz}$)

APCI-MASS : $m/z = 455$ ($M+H$)⁺

Preparation 330

Methyl 4-[5-(4-cyclohexylphenyl)-1,3,4-thiadiazol-2-yl]benzoate

IR (KBr) : 2925, 2850, 1716, 1432, 1274, 1108, 997 cm^{-1}

NMR (CDCl_3 , δ) : 1.2-1.6 (5H, m), 1.7-2.0 (5H, m),

2.58 (1H, m), 3.96 (3H, s), 7.34 (2H, d, J=8.2Hz),
7.93 (2H, d, J=8.2Hz), 8.07 (2H, d, J=8.6Hz), 8.16
(2H, d, J=8.6Hz)

APCI-MASS : m/z = 379 (M+H⁺)

5

Preparation 331

Methyl 4-[5-[4-(piperidin-1-yl)phenyl]-1,3,4-thiadiazol-2-yl]benzoate

10 IR (KBr) : 2940, 2848, 1720, 1602, 1436, 1415, 1276,
1108 cm⁻¹

NMR (CDCl₃, δ) : 1.68 (6H, br), 3.34 (4H, br), 3.96
(3H, s), 6.95 (2H, d, J=8.7Hz), 7.88 (2H, d,
J=8.7Hz), 8.05 (2H, d, J=8.6Hz), 8.16 (2H, d,
J=8.6Hz)

15 APCI-MASS : m/z = 380 (M+H⁺)

Preparation 332

Methyl 4-[5-(4-n-octyloxyphenyl)-1,3,4-thiadiazol-2-yl]benzoate

20 IR (KBr) : 2927, 2858, 1720, 1606, 1434, 1276,
1106 cm⁻¹

NMR (CDCl₃, δ) : 0.89 (3H, t, J=6.8Hz), 1.2-1.6 (10H,
m), 1.7-1.9 (2H, m), 3.96 (3H, s), 4.03 (2H, t,
J=6.5Hz), 7.00 (2H, d, J=8.9Hz), 7.95 (2H, d,
25 J=8.9Hz), 8.06 (2H, d, J=8.4Hz), 8.16 (2H, d,
J=8.4Hz)

APCI-MASS : m/z = 425 (M+H⁺)

Preparation 333

30 Methyl 4-[5-(4-trans-n-pentylcyclohexyl)-1,3,4-
thiadiazol-2-yl]benzoate

IR (KBr) : 2923, 2850, 1722, 1440, 1276, 1110 cm⁻¹

NMR (CDCl₃, δ) : 0.89 (3H, t, J=6.9Hz), 1.0-1.8 (13H,
m), 1.92 (2H, d, J=13.4Hz), 2.24 (2H, d, J=12.2Hz),
35 3.15 (1H, tt, J=12.2 and 3.5Hz), 3.95 (3H, s), 8.01

(2H, dd, J=8.6 and 2.0Hz), 8.13 (2H, dd, J=8.6 and 2.0Hz)

APCI-MASS : $m/z = 373$ ($M+H^+$)

5 Preparation 334

Methyl 4-[5-[4-(4-n-propyloxyphenyl)phenyl]-1,3,4-thiadiazol-2-yl]benzoate

IR (KBr) : 1720, 1540, 1508, 1282 cm^{-1}

10 NMR (CDCl_3 , δ) : 1.07 (3H, t, J=7.5Hz), 1.85 (2H, m), 3.9-4.1 (5H, m), 7.01 (2H, d, J=8.8Hz), 7.59 (2H, d, J=8.8Hz), 7.70 (2H, d, J=8.4Hz), 8.07 (2H, d, J=8.4Hz), 8.1-8.2 (4H, m)

APCI-MASS : $m/z = 431$ ($M+H$)⁺

15 Preparation 335

To a suspension of 4-hexyloxybenzoic acid in oxalyl chloride (10 ml) and dichloromethane (10 ml) was added N,N-dimethylformamide (0.1 ml). The mixture was stirred at room temperature for 2 hours. The solvent was removed under reduced pressure to give crude 4-hexyloxybenzoyl chloride. To a suspension of Ethyl 3-amino-4-hydroxybenzoate (733 mg) and triethylamine (1.38 ml) and 4-dimethylaminopyridine (DMAP, 10 mg) in methylene chloride (10 ml) was added the solution of 4-hexyloxybenzoyl chloride obtained above in dichloromethane (5 ml) dropwise at 10°C. The reaction mixture was stirred at 10°C for 1.5 hours and diluted with dichloromethane (20 ml). The solution was washed with H_2O (20 ml), 1N HCl aq. (20 ml x 2), H_2O (20 ml) and brine (20 ml) successively. The organic layer was dried over MgSO_4 and the solvent was removed under reduced pressure. To the residue was added toluene (15 ml) and p-toluenesulfonic acid (10 mg). The mixture was refluxed for 6 hours and the solvent was removed under reduced pressure. The residue was triturated with acetonitrile, and precipitate was collected with filtration and dried over P_2O_5 to give 2-(4-

20
25
30
35

Hexyloxyphenyl)-5-ethoxycarbonylbenzoxazole (0.60 g).

IR (KBr) : 2952, 2871, 1712, 1623, 1500, 1294,
1255 cm^{-1}

5 NMR (CDCl_3 , δ) : 0.92 (3H, t, $J=6.6\text{Hz}$), 1.3-1.6 (9H,
m), 1.7-1.9 (2H, m), 4.05 (2H, t, $J=6.5\text{Hz}$), 4.42
(2H, q, $J=7.1\text{Hz}$), 7.03 (2H, d, $J=6.9\text{Hz}$), 7.57 (1H,
d, $J=8.6\text{Hz}$), 8.08 (1H, dd, $J=8.6$ and 1.7Hz), 8.18
(2H, d, $J=6.9\text{Hz}$), 8.43 (1H, d, $J=1.7\text{Hz}$)

APCI-MASS : $m/z = 368$ ($\text{M}+\text{H}^+$)

10

The following compounds (Preparations 336 to 337) were
obtained according to a similar manner to that of Preparation
335.

15 Preparation 336

5-Ethoxycarbonyl-2-(2-octyloxypyridin-5-yl)benzoxazole

IR (KBr) : 2933, 2858, 1716, 1623, 1604, 1577, 1467,
1290, 1213, 1083 cm^{-1}

20 NMR (CDCl_3 , δ) : 0.89 (3H, t, $J=6.7\text{Hz}$), 1.2-1.5 (10H,
m), 1.43 (3H, t, $J=7.1\text{Hz}$), 1.7-1.9 (2H, m), 4.3-4.5
(4H, m), 6.87 (1H, d, $J=8.7\text{Hz}$), 7.60 (1H, d,
 $J=8.6\text{Hz}$), 8.11 (1H, dd, $J=8.6$ and 1.6Hz), 8.37 (1H,
dd, $J=8.8$ and 2.4Hz), 8.45 (1H, d, $J=1.6\text{Hz}$), 9.03
(1H, d, $J=2.4\text{Hz}$)

25 APCI-MASS : $m/z = 397$ ($\text{M}+\text{H}^+$)

Preparation 337

2-[4-(4-Hexylphenyl)phenyl]-5-ethoxycarbonylbenzoxazole

30 IR (KBr) : 2952, 2871, 1712, 1623, 1500, 1294, 1255,
1024 cm^{-1}

35 NMR (CDCl_3 , δ) : 0.90 (3H, t, $J=6.6\text{Hz}$), 1.2-1.5 (6H,
m), 1.44 (3H, t, $J=7.1\text{Hz}$), 1.6-1.8 (2H, m), 2.67
(2H, t, $J=7.3\text{Hz}$), 4.43 (2H, q, $J=7.1\text{Hz}$), 7.27 (1H,
d, $J=3.7\text{Hz}$), 7.32 (1H, s), 7.5-7.7 (3H, m), 7.77
(2H, d, $J=8.6\text{Hz}$), 8.12 (1H, dd, $J=8.6$ and 1.7Hz),

8.32 (2H, d, J=8.5Hz), 8.48 (1H, d, J=1.2Hz)

APCI-MASS : $m/z = 428$ ($M+H^+$)

Preparation 338

5 A suspension of 4-[4-(8-bromooctyloxy)phenyl]benzoic acid (1 g) in 2,6-dimethylmorpholine (3.06 ml) was refluxed for 30 minutes. The reaction mixture was added to a mixture of water and ethyl acetate and adjusted to pH 2.0 with conc. HCl. The organic layer was taken and dried over magnesium sulfate. The magnesium sulfate was filtered off, and the filtrate was evaporated under reduced pressure to give 4-[4-[8-(2,6-dimethylmorpholin-4-yl)octyloxy]phenyl]benzoic acid hydrochloride (0.95 g).

IR (KBr) : 2939.0, 1704.8, 1606.4, 1189.9 cm^{-1}

15 NMR (DMSO- d_6 , δ) : 1.12 (6H, d, J=6.3Hz), 1.2-1.6 (10H, m), 1.6-1.9 (4H, m), 2.4-2.7 (2H, m), 2.9-3.1 (2H, m), 3.8-4.0 (2H, m), 4.02 (2H, t, J=6.3Hz), 7.04 (2H, d, J=8.8Hz), 7.68 (2H, d, J=8.8Hz), 7.75 (2H, d, J=8.4Hz), 7.99 (2H, d, J=8.4Hz)

20 APCI-MASS : $m/z = 440$ ($M+H^+$)

Preparation 339

Sodium hydride (60% suspension in mineral oil, 108 mg) was added to ethoxyethanol (10 ml), and the solution was stirred at 60°C for 20 minutes. To the solution was added Methyl 4-[4-(8-bromooctyloxy)phenyl]benzoate (1.26 g), and the reaction mixture was stirred at 70°C for 2 hours. To the reaction mixture was added 10% sodium hydroxide aqueous solution (2.4 ml), and the solution was stirred at 70°C for 1 hour. After cooling, the solution was adjusted to pH 2.0 with 1N hydrochloric acid. The precipitate was collected by filtration, and dried to give 4-[4-[8-(2-Ethoxyethoxy)octyloxy]phenyl]benzoic acid (1.13 g).

35 IR (KBr) : 2933, 2858, 1685, 1604, 1434, 1294, 1132 cm^{-1}

NMR (DMSO- d_6 , δ) : 1.09 (3H, t, $J=7.0\text{Hz}$), 1.2-1.9
(14H, m), 3.2-3.6 (6H, m), 4.01 (2H, d, $J=6.3\text{Hz}$),
7.04 (2H, d, $J=8.8\text{Hz}$), 7.67 (2H, d, $J=8.8\text{Hz}$), 7.74
(2H, d, $J=8.5\text{Hz}$), 7.98 (2H, d, $J=8.5\text{Hz}$)

5 APCI-MASS : $m/z = 415$ ($M+H^+$)

The following compound was obtained according to a
similar manner to that of Preparation 300.

10 Preparation 340

4-n-Pentyloxybenzoylhydrazine

IR (KBr) : 3182, 2937, 2869, 1645, 1618, 1571,
1251 cm^{-1}

15 NMR (DMSO- d_6 , δ) : 0.89 (3H, d, $J=7.1\text{Hz}$), 1.2-1.8 (6H,
m), 4.00 (2H, t, $J=6.5\text{Hz}$), 4.41 (2H, s), 6.96 (2H,
d, $J=8.8\text{Hz}$), 7.78 (2H, d, $J=8.8\text{Hz}$), 9.59 (1H, s)

APCI-MASS : $m/z = 223$ ($M+H^+$)

20 The following compound was obtained according to a
similar manner to that of Preparation 291.

Preparation 341

1-(4-Methoxycarbonylbenzoyl)-2-(4-n-pentyloxybenzoyl)-
hydrazine

25 IR (KBr) : 3234, 2956, 2931, 1724, 1683, 1643, 1610,
1284, 1253 cm^{-1}

30 NMR (DMSO- d_6 , δ) : 0.90 (3H, t, $J=6.9\text{Hz}$), 1.2-1.5 (4H,
m), 1.6-1.8 (2H, m), 3.90 (3H, s), 4.04 (2H, t,
 $J=6.5\text{Hz}$), 7.04 (2H, d, $J=8.8\text{Hz}$), 7.90 (2H, d,
 $J=8.8\text{Hz}$), 8.03 (2H, d, $J=8.7\text{Hz}$), 8.10 (2H, d,
 $J=8.7\text{Hz}$), 10.42 (1H, s), 10.64 (1H, s)

APCI-MASS : $m/z = 385$ ($M+H^+$)

35 The following compound was obtained according to a
similar manner to that of Preparation 328.

Preparation 342

Methyl 4-[5-(4-n-pentyloxyphenyl)thiadiazol-2-yl]benzoate

IR (KBr) : 2940, 2871, 1720, 1606, 1438, 1280 cm^{-1}

NMR (CDCl_3 , δ) : 0.95 (3H, t, $J=7.1\text{Hz}$), 1.3-1.6 (4H, m), 1.8-2.0 (2H, m), 3.96 (3H, s), 4.03 (2H, t, $J=6.5\text{Hz}$), 6.99 (2H, d, $J=8.8\text{Hz}$), 7.94 (2H, d, $J=8.8\text{Hz}$), 8.06 (2H, d, $J=8.7\text{Hz}$), 8.16 (2H, d, $J=8.7\text{Hz}$)

APCI-MASS : $m/z = 383$ ($\text{M}+\text{H}^+$)

The following compound was obtained according to a similar manner to that of Preparation 32

Preparation 343

4-[5-(4-n-Pentyloxyphenyl)thiadiazol-2-yl]benzoic acid

IR (KBr) : 2954, 2867, 1687, 1602, 1432, 1294, 1255 cm^{-1}

NMR ($\text{DMSO}-d_6$, δ) : 0.91 (3H, t, $J=7.0\text{Hz}$), 1.3-1.5 (4H, m), 1.7-1.9 (2H, m), 4.07 (2H, t, $J=6.7\text{Hz}$), 7.13 (2H, d, $J=8.8\text{Hz}$), 7.97 (2H, d, $J=8.8\text{Hz}$), 8.07 (4H, s)

APCI-MASS : $m/z = 369$ ($\text{M}+\text{H}^+$)

The following compound was obtained according to a similar manner to that of Preparation 49.

Preparation 344

1-[4-[5-(4-n-Pentyloxyphenyl)thiadiazol-2-yl]benzoyl]-benzotriazole 3-oxide

IR (KBr) : 2948, 2873, 1770, 1602, 1257, 1232 cm^{-1}

NMR (CDCl_3 , δ) : 0.95 (3H, t, $J=7.1\text{Hz}$), 1.3-1.6 (4H, m), 1.8-2.0 (2H, m), 4.04 (2H, t, $J=6.5\text{Hz}$), 7.01 (2H, d, $J=8.1\text{Hz}$), 7.4-7.7 (3H, m), 7.97 (2H, d, $J=8.1\text{Hz}$), 8.12 (1H, d, $J=8.2\text{Hz}$), 8.24 (2H, d,

J=8.0Hz), 8.40 (2H, d, J=8.0Hz)

APCI-MASS : m/z = 486 (M+H⁺)

Preparation 345

5 To a solution of 4-bromobenzaldehyde oxime chloride (647 mg) and 4-n-pentyloxy-phenylacetylene (650 mg) in tetrahydrofuran (7 ml) was added triethylamine (0.5 ml) in tetrahydrofuran (5 ml) dropwise at 40°C. The solution was stirred at 40°C for 30 minutes, poured into water and
10 extracted with ethyl acetate. The organic layer was washed with H₂O, brine and dried over magnesium sulfate. The solvents were removed under reduced pressure and the residue was triturated with acetonitrile. The precipitate was collected by filtration and dried to give 4-[5-(4-n-pentyloxyphenyl)isoxazol-3-yl]bromobenzene (0.59 g).

15 IR (KBr) : 2948, 2867, 1612, 1430, 1255 cm⁻¹

NMR (CDCl₃, δ) : 0.95 (3H, t, J=6.9Hz), 1.3-1.6 (4H, m), 1.7-1.9 (2H, m), 4.01 (2H, t, J=6.5Hz), 6.66 (1H, s), 6.98 (2H, d, J=8.8Hz), 7.60 (2H, d, J=8.6Hz), 7.7-7.9 (4H, m)

20 APCI-MASS : m/z = 388 (M+H⁺)

Preparation 346

25 To a suspension of 4-[5-(4-n-pentyloxyphenyl)isoxazol-3-yl]bromobenzene (386 mg) in tetrahydrofuran (5 ml) was added 1.55M n-butyllithium in hexane (0.84 ml) at -40°C under N₂ stream and the solution was stirred for 1 hour at -40°C. To the solution was added crushed dryice (1 g) and the suspension was stirred for 1 hour at -40°C. The suspension
30 was diluted with H₂O, and acidified with 1N-hydrochloric acid. The precipitate was collected by filtration and dried to give 4-[5-(4-n-pentyloxyphenyl)isoxazol-3-yl]benzoic acid (312 mg).

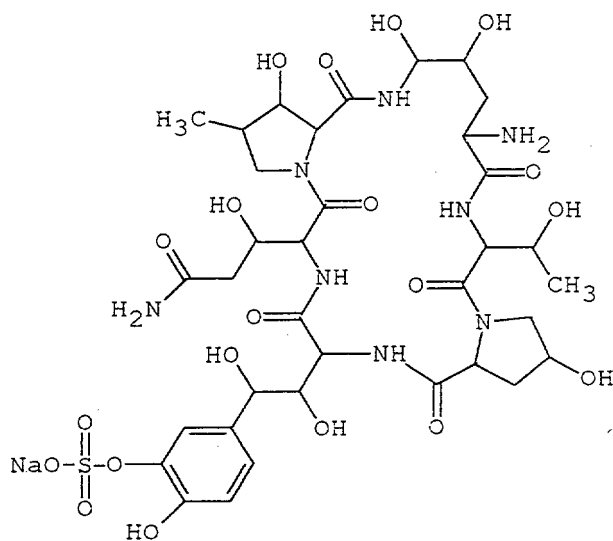
35 IR (KBr) : 2939, 2867, 1681, 1614, 1429, 1255, 1178, 821 cm⁻¹

NMR (DMSO-d₆, δ) : 0.91 (3H, t, J=7.1Hz), 1.3-1.5 (4H, m), 1.6-1.8 (2H, m), 4.04 (2H, t, J=6.5Hz), 7.11 (2H, d, J=8.9Hz), 7.54 (1H, s), 7.85 (2H, d, J=8.9Hz), 7.98 (2H, d, J=8.6Hz), 8.11 (2H, d, J=8.6Hz)

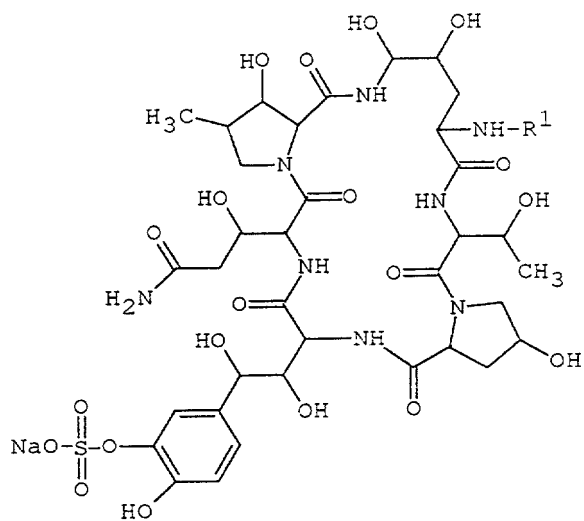
APCI-MASS : m/z = 352 (M+H⁺)

The Starting Compound in the following Examples 1 to 117 and The Object Compounds (1) to (122) and (124) in the following Examples 1 to 122 and 124 are illustrated by chemical formulae as below.

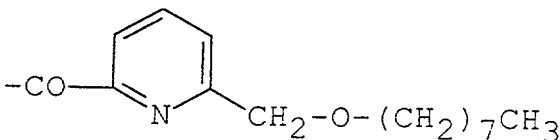
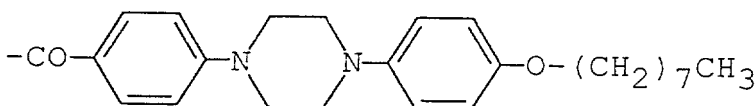
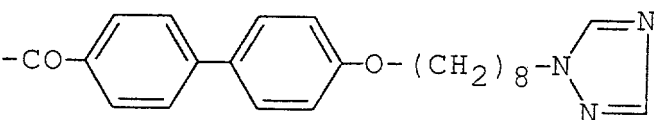
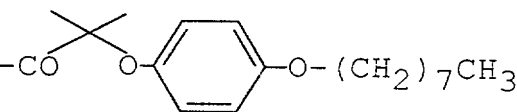
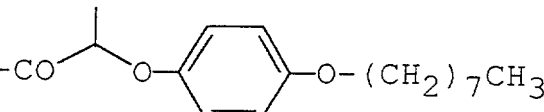
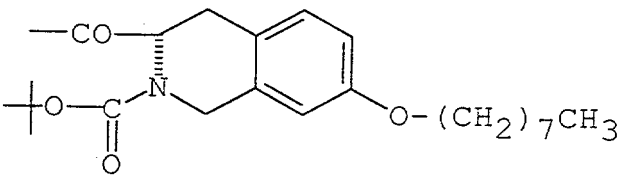
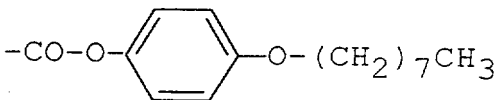
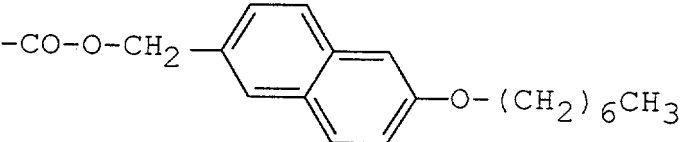
The Starting Compound
(the same in
Examples 1 to 117)



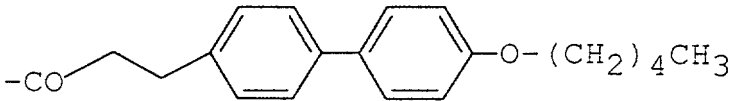
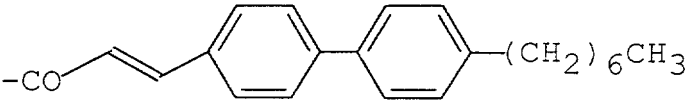
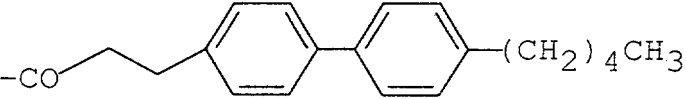
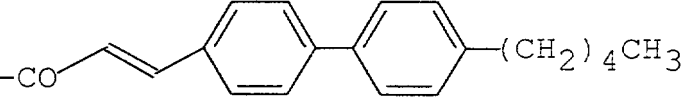
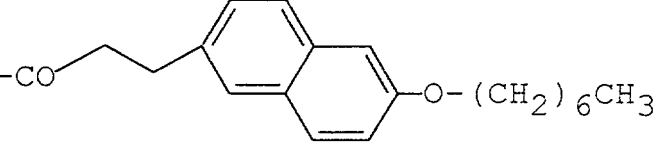
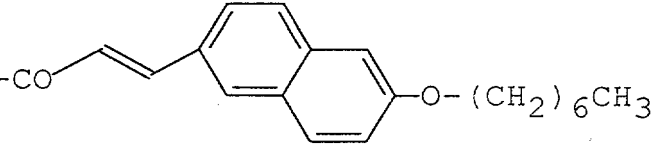
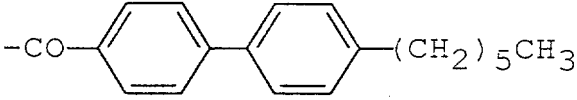
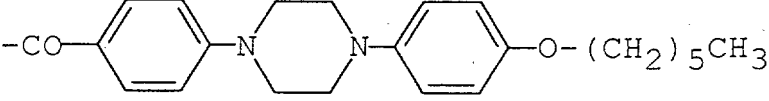
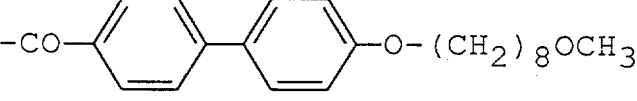
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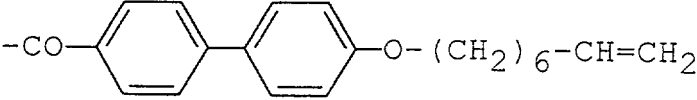
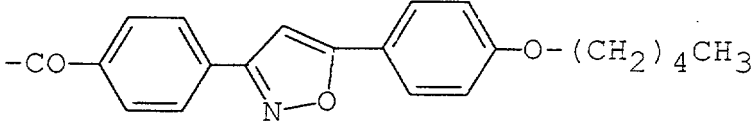
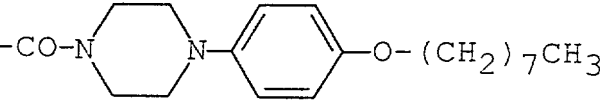
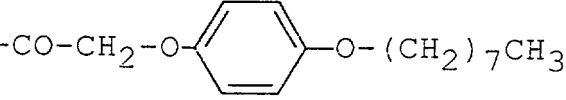
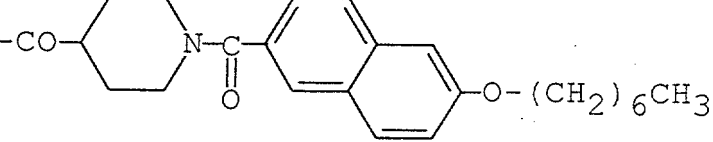
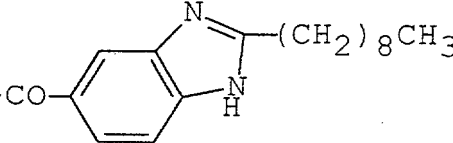
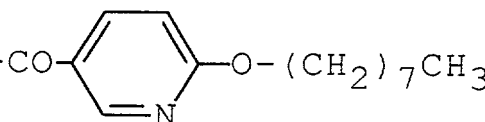
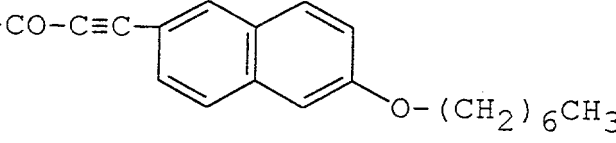


In the following Examples, The Object Compound (X) [e.g. The Object Compound (1)] means the object compound of Example (X) [e.g. Example (1)].

Example No.	R^1
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Example No.	R ¹
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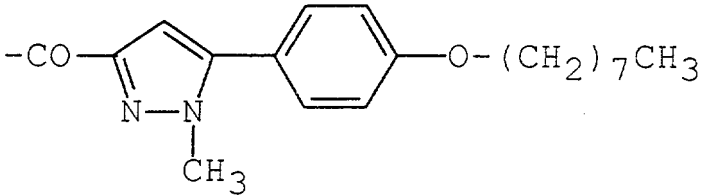
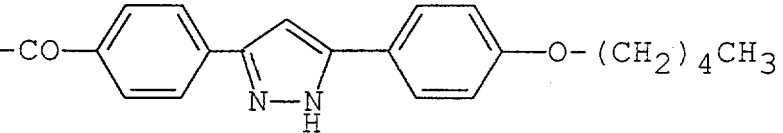
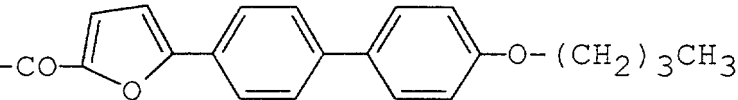
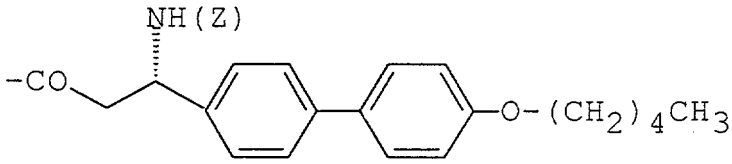
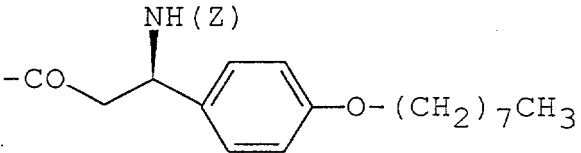
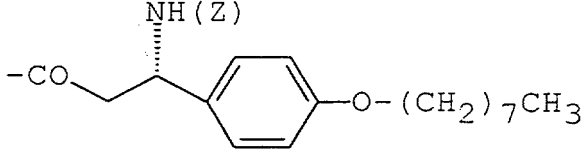
Example No.	R ¹
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23	
24 major product	

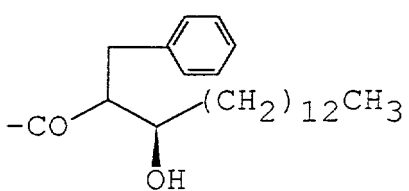
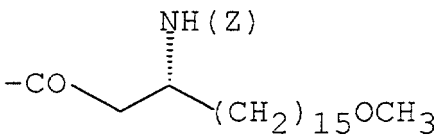
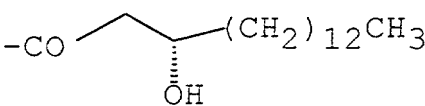
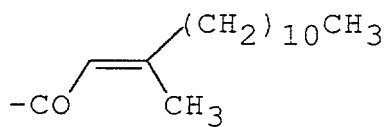
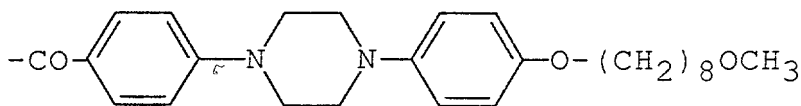
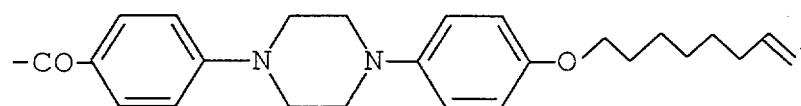
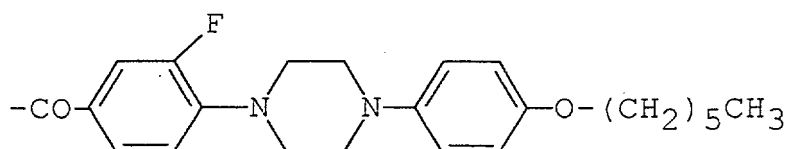
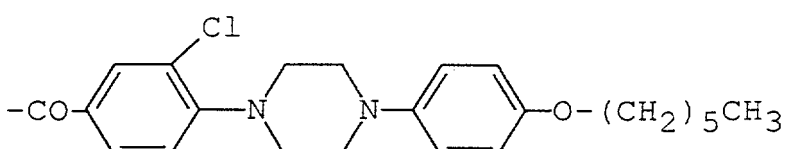
Example No.	R ¹
24 minor product	
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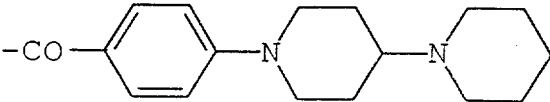
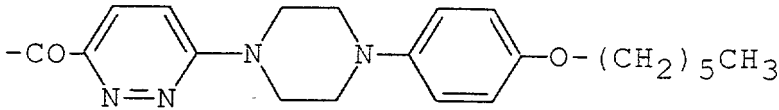
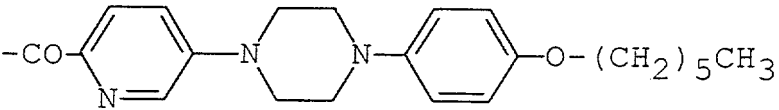
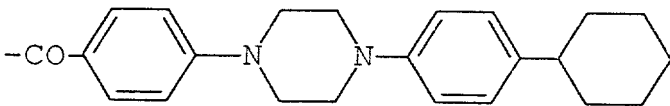
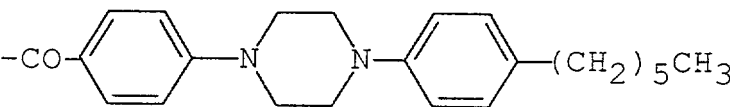
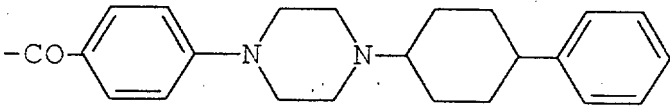
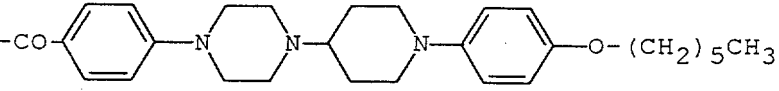
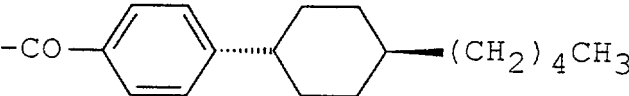
Example No.	R ¹
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38 major product	
38 minor product	

Example No.	R ¹
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42 mixture product	
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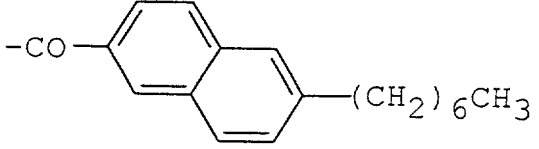
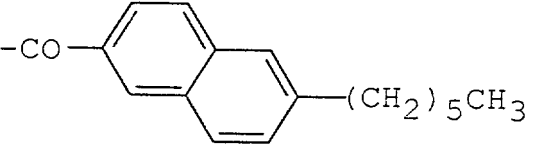
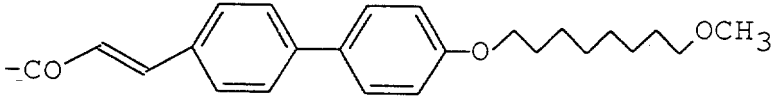
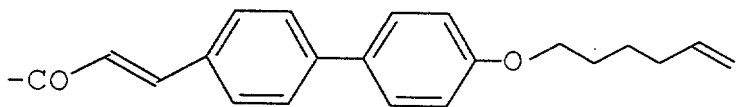
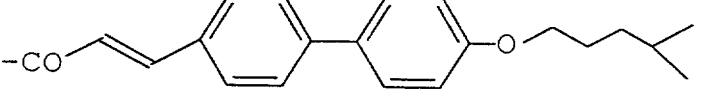
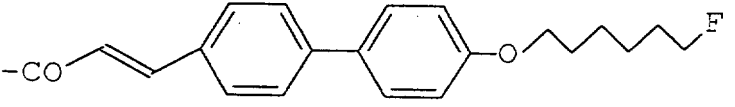
Example No.	R ¹
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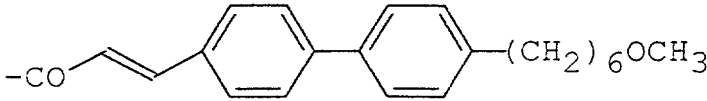
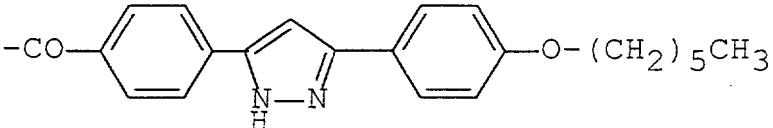
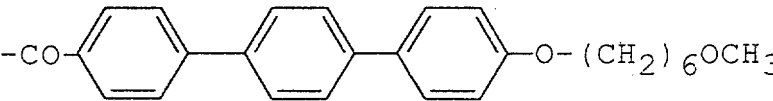
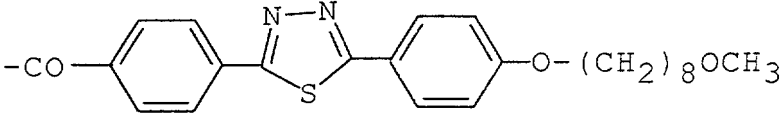
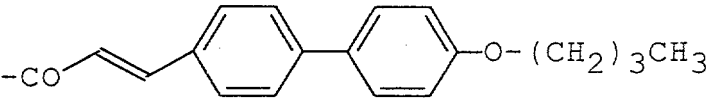
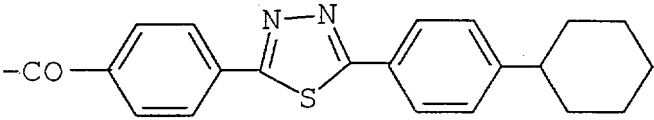
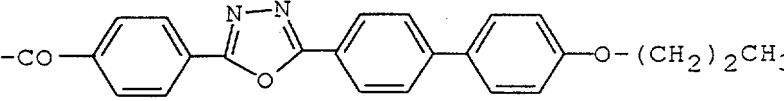
Example No.	R^1
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56	
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58	
59	

Example No.	R ¹
60	
61	
62	
63	
64 major product	
64 minor product	
65	
66	

Example No.	R ¹
67	
68	
69	
70	
71	
72	
73	
74	

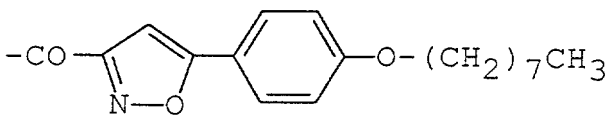
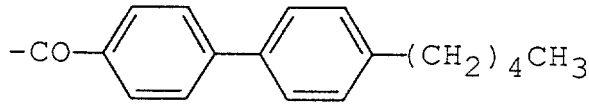
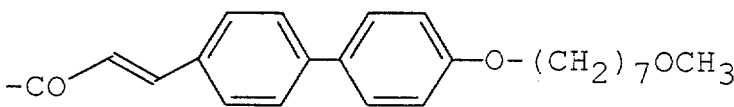
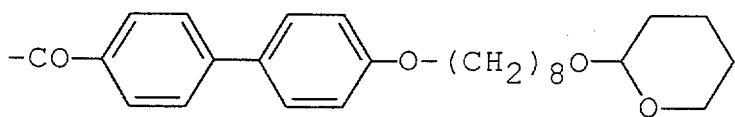
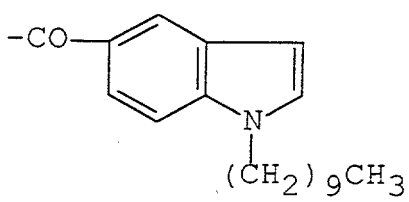
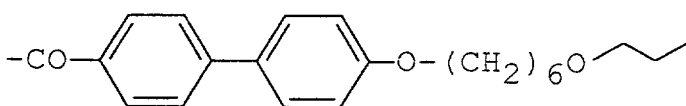
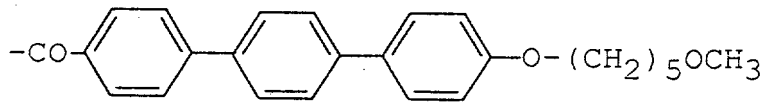
Example No.	R ¹
75	
76	
77	
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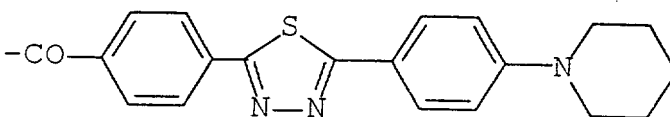
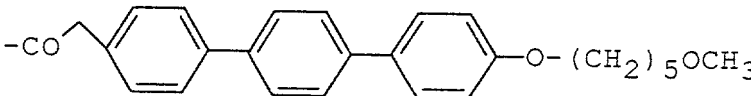
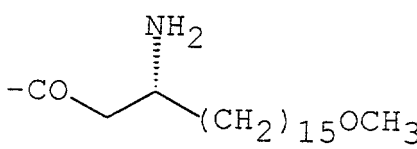
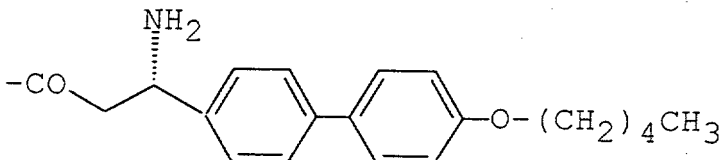
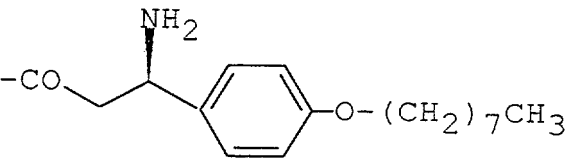
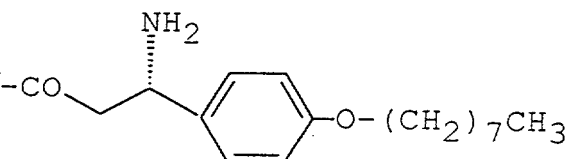
Example No.	R^1
81	
82	
83	
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85	
86	

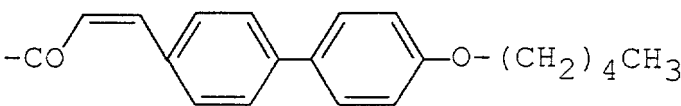
Example No.	R ¹
87	
88	
89	
90	
91	
92	
93	

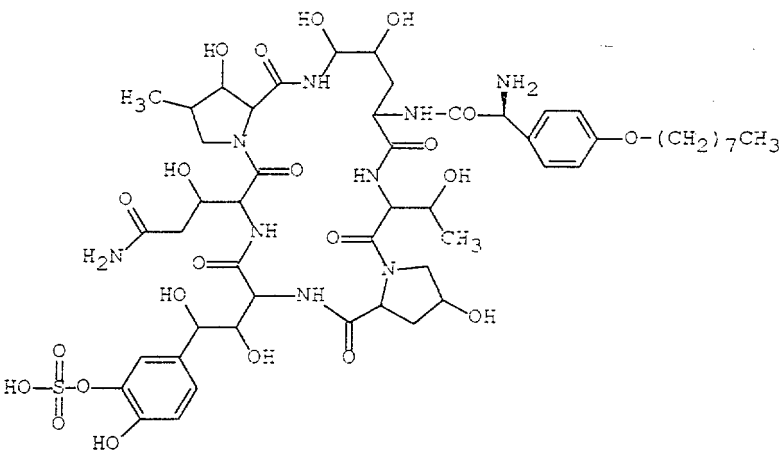
Example No.	R ¹
94	
95	
96	
97	
98	
99	
100	
101	

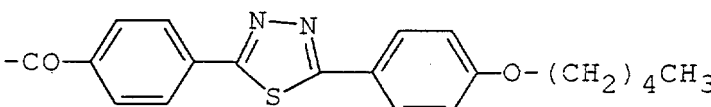
Example No.	R ¹
102	
103	
104	
105	
106	
107	
108	

Example No.	R ¹
109	
110	
111	
112	
113	
114	
115	

Example No.	R ¹
116	
117	
118	
119	
120	
121	

Example No.	R^1
122	

Example No.	The Object Compound
123	

Example No.	R^1
124	

Example 1

To a solution of The Starting Compound (1 g) and 1-(6-octyl-oxymethylpicolinoyl)benzotriazole 3-oxide (0.399 g) in N,N-dimethylformamide (10 ml) was added 4-(N,N-dimethylamino)pyridine (0.140 g), and stirred for 12 hours at ambient temperature. The reaction mixture was pulverized with ethyl acetate. The precipitate was collected by filtration, and dried under reduced pressure. The powder was dissolved in water, and subjected to column chromatography on ion exchange resin (DOWEX-50WX4 (Trademark : prepared by Dow Chemical)) eluting with water. The fractions containing the object compound were combined, and subjected to column chromatography on ODS (YMC-gel-ODS-AM-S-50) (Trademark : prepared by Yamamura Chemical Lab.) eluting with 50% methanol aqueous solution. The fractions containing the object compound were combined, and evaporated under reduced pressure to remove methanol. The residue was lyophilized to give The Object Compound (1).

IR (KBr) : 3347, 1664, 1629, 1517 cm^{-1}

NMR (DMSO- d_6 , δ) : 0.86 (3H, t, $J=6.7\text{Hz}$), 0.98 (3H, d, $J=6.7\text{Hz}$), 1.09 (3H, d, $J=6.0\text{Hz}$), 1.2-1.47 (10H, m), 1.47-1.67 (2H, m), 1.67-2.06 (3H, m), 2.06-2.5 (4H, m), 3.19 (1H, m), 3.53 (2H, t, $J=6.4\text{Hz}$), 3.5-3.85 (2H, m), 3.85-4.7 (13H, m), 5.35 (11H, m), 5.56 (1H, d, $J=5.7\text{Hz}$), 6.73 (1H, d, $J=8.3\text{Hz}$), 6.83 (1H, d, $J=8.3\text{Hz}$), 6.89 (1H, s), 7.05 (1H, s), 7.11 (1H, s), 7.32 (1H, m), 7.43 (1H, d, $J=8.5\text{Hz}$), 7.63 (1H, d, $J=7.3\text{Hz}$), 7.85-8.13 (4H, m), 8.66 (1H, d, $J=7.8\text{Hz}$), 8.84 (1H, s)

FAB-MASS : $m/z = 1228$ ($M^+ + \text{Na}$)

Elemental Analysis Calcd. for $\text{C}_{50}\text{H}_{72}\text{N}_9\text{O}_{22}\text{SNa} \cdot 6\text{H}_2\text{O}$:

C 45.49, H 6.44, N 9.59

Found : C 45.89, H 6.52, N 9.69

The Object Compounds (2) to (25) were obtained according to a similar manner to that of Example 1.

Example 2

5 IR (KBr) : 3353, 1666, 1510, 1236 cm^{-1}
NMR (DMSO- d_6 , δ) : 0.86 (3H, t, $J=6.7\text{Hz}$), 0.96 (3H, d, $J=6.7\text{Hz}$), 1.06 (3H, d, $J=5.8\text{Hz}$), 1.2-1.5 (10H, m), 1.55-2.05 (5H, m), 2.11-2.7 (4H, m), 3.0-3.3 (5H, m), 3.3-3.5 (4H, m), 3.6-4.5 (15H, m), 4.6-5.6 (12H, m), 6.6-7.2 (10H, m), 7.2-7.5 (3H, m), 7.81 (2H, d, $J=8.8\text{Hz}$), 8.05 (1H, d, $J=8.7\text{Hz}$), 8.28 (1H, d, $J=8.7\text{Hz}$), 8.41 (1H, d, $J=6.7\text{Hz}$), 8.84 (1H, s)
FAB-MASS : $m/z = 1373$ ($\text{M}^+ + \text{Na}$)
15 Elemental Analysis Calcd. for $\text{C}_{60}\text{H}_{83}\text{N}_{10}\text{O}_{22}\text{SNa} \cdot 4\text{H}_2\text{O}$:
C 50.63, H 6.44, N 9.84
Found : C 50.59, H 6.59, N 9.79

Example 3

20 IR (KBr) : 3350, 1664, 1627, 1047 cm^{-1}
NMR (DMSO- d_6 , δ) : 0.96 (3H, d, $J=6.6\text{Hz}$), 1.08 (3H, d, $J=5.7\text{Hz}$), 1.15-1.53 (8H, m), 1.55-2.1 (9H, m), 2.1-2.45 (3H, m), 2.5-2.7 (1H, m), 3.18 (1H, m), 3.6-3.83 (2H, m), 3.83-4.6 (17H, m), 4.7-5.4 (11H, m), 5.51 (1H, d, $J=5.9\text{Hz}$), 6.73 (1H, d, $J=8.2\text{Hz}$), 6.83 (1H, d, $J=8.2\text{Hz}$), 6.85 (1H, s), 7.03 (2H, d, $J=8.4\text{Hz}$), 7.05 (1H, s), 7.30 (1H, s), 7.2-7.5 (2H, m), 7.67 (2H, d, $J=8.4\text{Hz}$), 7.71 (2H, d, $J=7.4\text{Hz}$), 7.94 (1H, s), 7.96 (2H, d, $J=7.4\text{Hz}$), 8.06 (1H, d, $J=8.0\text{Hz}$), 8.25 (1H, d, $J=6.7\text{Hz}$), 8.50 (1H, s), 8.74 (1H, d, $J=6.7\text{Hz}$), 8.84 (1H, s)
25 FAB-MASS : $m/z = 1356$ ($\text{M}^+ + \text{Na}$)
Elemental Analysis Calcd. for $\text{C}_{58}\text{H}_{76}\text{N}_{11}\text{O}_{22}\text{SNa} \cdot 4\text{H}_2\text{O}$:
30 C 49.53, H 6.02, N 10.95
35

Found : C 49.26, H 6.22, N 10.77

Example 4

IR (KBr) : 3350, 1660, 1631, 1047 cm^{-1}

5 NMR (DMSO- d_6 , δ) : 0.86 (3H, t, $J=6.9\text{Hz}$), 0.97 (3H, d, $J=6.6\text{Hz}$), 1.09 (3H, d, $J=5.3\text{Hz}$), 1.2-1.5 (10H, m), 1.37 (6H, s), 1.55-2.0 (5H, m), 2.1-2.6 (4H, m), 3.16 (1H, m), 3.73 (2H, m), 3.89 (2H, t, $J=6.3\text{Hz}$), 3.95-4.49 (11H, m), 4.68-5.21 (10H, m), 5.25 (1H, d, $J=4.1\text{Hz}$), 5.53 (1H, d, $J=5.7\text{Hz}$), 6.73 (1H, d, $J=8.2\text{Hz}$), 6.75-6.85 (4H, m), 6.91 (1H, d, $J=8.2\text{Hz}$), 7.05 (1H, s), 7.15 (1H, s), 7.3-7.5 (2H, m), 7.9-8.2 (3H, m), 8.84 (1H, s)

15 FAB-MASS : $m/z = 1271$ ($\text{M}^+ + \text{Na}$)

Elemental Analysis Calcd. For $\text{C}_{53}\text{H}_{77}\text{N}_8\text{O}_{23}\text{SNa}\cdot 4\text{H}_2\text{O}$:

C 48.18, H 6.48, N 8.48

Found : C 48.04, H 6.51, N 8.38

20 Example 5

IR (KBr) : 1666, 1629, 1222 cm^{-1}

NMR (DMSO- d_6 , δ) : 0.85 (3H, t, $J=6.6\text{Hz}$), 0.9-1.12 (6H, m), 1.12-1.52 (13H, m), 1.52-1.93 (5H, m), 2.08-2.55 (4H, m), 3.16 (1H, m), 3.6-5.3 (26H, m), 5.49 + 5.54 (1H, d, $J=5.8\text{Hz}$, mixture of diastereomer), 6.60-7.1 (7H, m), 7.04 (1H, s), 7.1 (1H, m), 7.2-7.5 (2H, m), 7.9-8.43 (3H, m), 8.83 (1H, s)

FAB-MASS : $m/z = 1257$ ($\text{M}^+ + \text{Na}$)

30 Elemental Analysis Calcd. for $\text{C}_{52}\text{H}_{75}\text{N}_8\text{O}_{23}\text{SNa}\cdot 3\text{H}_2\text{O}$:

C 48.44, H 6.33, N 8.69

Found : C 48.16, H 6.51, N 8.53

Example 6

35 IR (KBr) : 3349, 1666, 1629, 1259 cm^{-1}

NMR (DMSO-d₆, δ) : 0.86 (3H, t, J=6.7Hz), 0.9 (3H, d, J=5.7Hz), 0.96 (3H, d, J=6.7Hz), 1.1-1.55 (19H, m), 1.55-2.0 (5H, m), 2.0-2.47 (4H, m), 2.65-3.25 (3H, m), 3.5-5.13 (27H, m), 5.17 (1H, d, J=3.2Hz), 5.24 (1H, d, J=4.5Hz), 5.38 (1H, d, J=5.9Hz), 6.5-6.9 (5H, m), 6.9-7.1 (3H, m), 7.2-7.46 (2H, m), 7.7-8.1 (3H, m), 8.83 (1H, s)

FAB-MASS : m/z = 1368 (M⁺+Na)

Elemental Analysis Calcd. for C₅₈H₈₄N₉O₂₄SNa·5H₂O :

C 48.50, N 6.60, N 8.78

Found : C 48.47, H 6.83, N 8.78

Example 7

IR (KBr) : 3350, 1666, 1502, 1199 cm⁻¹

NMR (DMSO-d₆, δ) : 0.86 (3H, t, J=6.6Hz), 0.97 (3H, d, J=6.7Hz), 1.06 (3H, d, J=5.7Hz), 1.2-1.5 (10H, m), 1.55-2.0 (5H, m), 2.1-2.6 (4H, m), 3.17 (1H, m), 3.7-4.5 (15H, m), 4.7-5.22 (10H, m), 5.24 (1H, d, J=4.4Hz), 5.60 (1H, d, J=5.9Hz), 6.68-7.03 (8H, m), 7.04 (1H, s), 7.2-7.42 (2H, m), 7.85-8.1 (3H, m), 8.83 (1H, s)

FAB-MASS : m/z = 1229 (M⁺+Na)

Elemental Analysis Calcd. for C₅₀H₇₁N₈O₂₃SNa·5H₂O :

C 46.29, H 6.29, N 8.64

Found : C 46.39, H 6.05, N 8.72

Example 8

IR (KBr) : 3350, 1666, 1631, 1513 cm⁻¹

NMR (DMSO-d₆, δ) : 0.88 (3H, t, J=6.2Hz), 0.97 (3H, d, J=6.7Hz), 1.04 (3H, d, J=5.7Hz), 1.2-1.58 (8H, m), 1.58-2.0 (5H, m), 2.0-2.6 (4H, m), 3.17 (1H, m), 3.6-4.5 (15H, m), 4.63-5.33 (13H, m), 5.53 (1H, d, J=5.9Hz), 6.73 (1H, d, J=8.2Hz), 6.82 (1H, d, J=8.2Hz), 6.84 (1H, s), 6.95-7.52 (7H, m), 7.66 (1H, d, J=7.6Hz), 7.7-7.9 (3H, m),

8.05 (1H, d, J=9.1Hz), 8.15 (1H, d, J=7.6Hz),
8.85 (1H, s)

FAB-MASS : $m/z = 1279$ ($M^+ + Na$)

Elemental Analysis Calcd. for $C_{54}H_{73}N_8O_{23}SNa \cdot 5H_2O$:

C 48.14, H 6.21, N 8.32

Found : C 48.43, H 6.28, N 8.30

Example 9

IR (KBr) : 3347, 2956, 1664, 1633, 1508, 1444, 1268,
1047 cm^{-1}

NMR (DMSO- d_6 , δ) : 0.9-1.1 (9H, m), 1.06 (3H, d,
J=5.9Hz), 1.3-1.5 (8H, m), 1.6-2.0 (7H, m), 2.1-
2.4 (3H, m), 2.5-2.6 (1H, m), 3.1-3.3 (1H, m),
3.6-4.4 (17H, m), 4.7-5.0 (8H, m), 5.09 (1H, d,
J=5.5Hz), 5.16 (1H, d, J=3.1Hz), 5.24 (1H, d,
J=4.5Hz), 6.73 (1H, d, J=8.2Hz), 6.8-6.9 (2H,
m), 6.98 (1H, d, J=8.3Hz), 7.05 (1H, d,
J=1.7Hz), 7.3-7.6 (5H, m), 8.08 (1H, d,
J=8.9Hz), 8.25 (1H, d, J=8.4Hz), 8.54 (1H, d,
J=7.5Hz), 8.83 (1H, s)

FAB-MASS : $m/z = 1257$ ($M^+ + Na$)

Elemental Analysis Calcd. for $C_{52}H_{75}N_8O_{23}SNa \cdot 4H_2O$:

C 47.78, H 6.40, N 8.57

Found : C 47.88, H 6.71, N 8.53

Example 10

IR (KBr) : 3350, 2931, 1664, 1625, 1529, 1440, 1276,
1226, 1047 cm^{-1}

NMR (DMSO- d_6 , δ) : 0.86 (3H, t, J=6.8Hz), 0.97 (3H,
d, J=6.7Hz), 1.12 (3H, d, J=5.9Hz), 1.2-1.5
(10H, m), 1.6-2.1 (5H, m), 2.1-2.4 (4H, m), 3.1-
3.3 (1H, m), 3.5-4.6 (15H, m), 4.7-5.0 (3H, m),
5.0-5.2 (7H, m), 5.27 (1H, d, J=4.4Hz), 5.55
(1H, d, J=5.7Hz), 6.73 (1H, d, J=8.2Hz), 6.8-7.0
(2H, m), 7.0-7.2 (4H, m), 7.3-7.6 (2H, m), 7.90

(1H, d, J=8.8Hz), 8.0-8.2 (2H, m), 8.8-8.9 (2H, m), 9.06 (1H, d, J=7.2Hz)

FAB-MASS : m/z = 1281 (M⁺+Na)

Elemental Analysis Calcd. for C₅₃H₇₁N₈O₂₄SNa·5H₂O :

5 C 47.18, H 6.05, N 8.30

Found : C 46.97, H 6.27, N 8.22

Example 11

10 NMR (DMSO-d₆, δ) : 0.87-1.05 (6H, m), 1.10 (3H, d, J=5.7Hz), 1.3-1.5 (4H, m), 1.6-1.9 (5H, m), 2.2-2.5 (3H, m), 2.6 (1H, m), 3.1-3.2 (1H, m), 3.7-4.5 (15H, m), 4.8-5.1 (8H, m), 5.09 (1H, d, J=5.64Hz), 5.16 (1H, d, J=3.2Hz), 5.26 (1H, d, J=4.2Hz), 5.52 (1H, d, J=6.0Hz), 6.73 (2H, d, J=8.4Hz), 6.8-6.9 (2H, m), 7.0-7.1 (3H, m), 7.2-7.4 (4H, m), 7.6-7.8 (6H, m), 8.11 (1H, d, J=8.4Hz), 8.29 (1H, d, J=8.4Hz), 8.51 (1H, d, J=7.7Hz), 8.85 (1H, s)

15 FAB-MASS : m/z = 1273 (M⁺+Na)

20 Elemental Analysis Calcd. for C₅₅H₇₁N₈O₂₂SNa·4H₂O :

C 49.92, H 6.02, N 8.47

Found : C 49.79, H 6.14, N 8.45

Example 12

25 IR (KBr) : 3330, 2929, 1670, 1629, 1533, 1440, 1280, 1226, 1045, 804 cm⁻¹

30 NMR (DMSO-d₆, δ) : 0.86 (3H, t, J=6.7Hz), 0.97 (3H, d, J=6.7Hz), 1.08 (3H, d, J=5.9Hz), 1.2-1.6 (10H, m), 1.6-2.0 (5H, m), 2.1-2.5 (4H, m), 3.1-3.3 (1H, m), 3.6-4.5 (15H, m), 4.8-5.1 (9H, m), 5.17 (1H, d, J=3.0Hz), 5.25 (1H, d, J=4.5Hz), 5.56 (1H, d, J=5.6Hz), 6.73 (1H, d, J=8.2Hz), 6.83 (1H, d, J=6.8Hz), 7.1-7.2 (3H, m), 7.3-7.5 (3H, m), 7.85 (1H, d, J=8.8Hz), 8.0-8.2 (3H, m), 8.84 (1H, s), 8.96 (1H, d, J=7.2Hz)

FAB-MASS : $m/z = 1269$ ($M^+ + Na$)

Elemental Analysis Calcd. for $C_{52}H_{71}N_8O_{22}S_2Na \cdot 4H_2O$:

C 47.34, H 6.04, N 8.49

Found : C 47.21, H 5.96, N 8.41

5

Example 13

IR (KBr) : 3345, 2927, 1664, 1629, 1515, 1442,
1274, 1047 cm^{-1}

10 NMR (DMSO- d_6 , δ) : 0.85 (3H, t, $J=6.7Hz$), 0.97 (3H,
d, $J=6.7Hz$), 1.10 (3H, d, $J=5.9Hz$), 1.2-1.4
(10H, m), 1.5-2.5 (8H, m), 2.46 (3H, s), 2.69
(2H, t, $J=7.7Hz$), 3.1-3.4 (2H, m), 3.6-4.5 (17H,
m), 4.8-5.2 (8H, m), 6.7-7.0 (3H, m), 7.05 (1H,
15 d, $J=1.7Hz$), 7.14 (1H, s), 7.3-7.6 (5H, m), 8.0-
8.2 (2H, m), 8.47 (1H, d, $J=7.0Hz$), 8.84 (1H, s)

FAB-MASS : $m/z = 1251$ ($M^+ + Na$)

Elemental Analysis Calcd. for $C_{53}H_{73}N_8O_{22}SNa \cdot 3H_2O$:

C 49.61, H 6.21, N 8.73

Found : C 49.88, H 6.44, N 8.74

20

Example 14

IR (KBr) : 3340, 1672, 1627, 1542, 1513, 1440, 1268,
1045 cm^{-1}

25 NMR (DMSO- d_6 , δ) : 0.84 (3H, t, $J=6.7Hz$), 0.94 (3H,
d, $J=6.7Hz$), 1.07 (3H, d, $J=6.0Hz$), 1.2-1.4
(12H, m), 1.6-2.0 (5H, m), 2.1-2.4 (3H, m), 2.6
(1H, m), 2.96 (2H, t, $J=7.4Hz$), 3.1-3.3 (1H, m),
3.6-4.5 (13H, m), 4.7-5.2 (11H, m), 5.50 (1H, d,
30 $J=5.7Hz$), 6.73 (1H, d, $J=8.2Hz$), 6.8-6.9 (2H,
m), 7.04 (1H, s), 7.2-7.5 (3H, m), 7.72 (1H, d,
 $J=8.5Hz$), 7.91 (1H, d, $J=8.4Hz$), 8.05 (1H, d,
 $J=8.4Hz$), 8.2-8.4 (1H, m), 8.80 (1H, d,
35 $J=7.7Hz$), 8.83 (1H, s)

FAB-MASS : $m/z = 1252$ ($M^+ + Na$)

Elemental Analysis Calcd. for $C_{52}H_{72}N_9O_{22}SNa \cdot 6H_2O$:

35

C 46.67, H 6.33, N 9.42

Found : C 46.72, H 6.53, N 9.45

Example 15

5 IR (KBr) : 3350, 2935, 1664, 1627, 1517, 1446, 1251,
1045 cm^{-1}

NMR (DMSO- d_6 , δ) : 0.90-1.1 (6H, m), 1.10 (3H, d,
J=5.9Hz), 1.2-1.4 (6H, m), 1.6-2.4 (8H, m), 2.6-
2.7 (1H, m), 3.1-3.3 (1H, m), 3.7-4.5 (16H, m),
10 4.7-5.4 (11H, m), 5.51 (1H, d, J=5.6Hz), 6.7-7.0
(3H, m), 7.0-7.6 (7H, m), 7.74 (1H, d, J=8.6Hz),
8.0-8.4 (5H, m), 8.7-8.8 (1H, m), 8.84 (1H, s)

FAB-MASS : m/z = 1301 (M^+ +Na)

Elemental Analysis Calcd. for $\text{C}_{55}\text{H}_{71}\text{N}_{10}\text{O}_{22}\text{SNa}\cdot 6\text{H}_2\text{O}$:

15 C 47.62, H 6.03, N 10.01

Found : C 47.65, H 6.03, N 10.03

Example 16

IR (Nujol) : 3353, 1668, 1627, 1540, 1515, 1500 cm^{-1}

20 NMR (DMSO- d_6 , δ) : 0.80-1.00 (6H, m), 1.06 (3H, d,
J=5.9Hz), 1.20-1.53 (4H, m), 1.60-1.95 (5H, m),
2.00-2.65 (8H, m), 2.80 (2H, t, J=7.5Hz), 3.05-
3.45 (1H, m), 3.50-3.85 (2H, m), 3.90-4.48 (11H,
m), 4.65-5.38 (11H, m), 5.47 (1H, d, J=6.0Hz),
25 6.65-6.90 (2H, m), 6.90-7.10 (2H, m), 7.10-7.65
(11H, m), 7.90-8.25 (2H, m), 8.30 (1H, d,
J=7.8Hz), 8.84 (1H, s)

FAB-MASS : m/z = 1275.3 (M^+ +Na)

Elemental Analysis Calcd. for $\text{C}_{55}\text{H}_{73}\text{N}_8\text{O}_{22}\text{SNa}\cdot 3\text{H}_2\text{O}$:

30 C 50.53, H 6.09, N 8.57

Found : C 50.48, H 6.39, N 8.57

Example 17

IR (Nujol) : 3351, 1656, 1623, 1538, 1515 cm^{-1}

35 NMR (DMSO- d_6 , δ) : 0.86 (3H, t, J=6.7Hz), 0.96 (3H,

d, J=6.7Hz), 1.08 (3H, d, J=5.8Hz), 1.15-1.40
(8H, m), 1.50-2.00 (5H, m), 2.10-2.48 (4H, m),
2.52-2.70 (2H, m), 3.05-3.28 (1H, m), 3.60-4.50
(13H, m), 4.70-5.20 (9H, m), 5.25 (1H, d,
5 J=4.6Hz), 5.52 (1H, d, J=6.0Hz), 6.68-6.92 (4H,
m), 7.04 (1H, d, J=1.0Hz), 7.22-7.50 (5H, m),
7.55-7.82 (7H, m), 8.14 (1H, d, J=8.4Hz), 8.31
(1H, d, J=8.4Hz), 8.54 (1H, d, J=7.7Hz), 8.84
(1H, s)

10 FAB-MASS : m/z = 1285 (M⁺+Na)

Example 18

IR (Nujol) : 3351, 1668, 1627, 1540, 1515 cm⁻¹

15 NMR (DMSO-d₆, δ) : 0.87 (3H, t, J=6.8Hz), 0.96 (3H,
d, J=6.7Hz), 1.06 (3H, d, J=5.8Hz), 1.17-1.48
(4H, m), 1.50-1.95 (5H, m), 2.05-2.70 (8H, m),
2.70-2.95 (2H, m), 3.05-3.30 (1H, m), 3.60-3.90
(2H, m), 3.90-4.50 (11H, m), 4.65-5.10 (9H, m),
5.15 (1H, d, J=3.2Hz), 5.23 (1H, d, J=4.2Hz),
20 5.48 (1H, d, J=6.0Hz), 6.67-6.90 (3H, m), 7.03
(1H, d, J=1.5Hz), 7.15-7.80 (11H, m), 8.00-8.20
(2H, m), 8.29 (1H, d, J=7.8Hz), 8.84 (1H, s)

FAB-MASS : m/z = 1259 (M⁺+Na)

Elemental Analysis Calcd. for C₅₅H₇₃N₈O₂₁SNa·6H₂O :

25 C 50.30, H 6.52, N 8.53

Found : C 50.42, H 6.50, N 8.45

Example 19

IR (Nujol) : 3351, 1668, 1652, 1623, 1540 cm⁻¹

30 NMR (DMSO-d₆, δ) : 0.87 (3H, t, J=6.7Hz), 0.96 (3H,
d, J=6.7Hz), 1.07 (3H, d, J=6.0Hz), 1.25-1.45
(4H, m), 1.50-2.00 (5H, m), 2.05-2.48 (4H, m),
2.50-2.75 (2H, m), 3.60-4.50 (13H, m), 4.68-5.25
(10H, m), 5.27 (1H, d, J=4.5Hz), 5.53 (1H, d,
35 J=6.0Hz), 6.67-6.98 (4H, m), 7.05 (1H, d,

J=1.0Hz), 7.22-7.58 (5H, m), 7.58-7.90 (7H, m),
8.16 (1H, d, J=9.0Hz), 8.34 (1H, d, J=8.4Hz),
8.57 (1H, d, J=7.7Hz), 8.85 (1H, s)

FAB-MASS : $m/z = 1258 (M^+ + Na)$

5 Elemental Analysis Calcd. for $C_{55}H_{71}N_8O_{21}SNa \cdot 5H_2O$:

C 49.84, H 6.15, N 8.45

Found : C 49.77, H 6.27, N 8.39

Example 20

10 IR (Nujol) : 3353, 1670, 1629, 1540, 1508 cm^{-1}
NMR (DMSO- d_6 , δ) : 0.88 (3H, t, J=6.5Hz), 0.97 (3H,
d, J=6.8Hz), 1.04 (3H, d, J=5.9Hz), 1.20-1.58
(8H, m), 1.60-1.96 (5H, m), 2.08-2.60 (6H, m),
2.70-3.00 (2H, m), 3.00-3.40 (1H, m), 3.60-3.85
15 (2H, m), 3.85-4.50 (13H, m), 4.50-5.60 (12H, m),
6.65-6.90 (3H, m), 7.00-7.15 (3H, m), 7.18-7.50
(4H, m), 7.59 (1H, s), 7.62-7.78 (2H, m), 7.95-
8.20 (2H, m), 8.30 (1H, d, J=7.7Hz), 8.83 (1H,
s)

20 FAB-MASS : $m/z = 1277 (M^+ + Na)$

Elemental Analysis Calcd. for $C_{55}H_{75}N_8O_{22}SNa \cdot 4H_2O$:

C 49.77, H 6.30, N 8.44

Found : C 49.67, H 6.31, N 8.40

25 Example 21

IR (Nujol) : 3351, 1654, 1623, 1538, 1515 cm^{-1}

NMR (DMSO- d_6 , δ) : 0.87 (3H, t, J=6.7Hz), 0.97 (3H,
d, J=6.7Hz), 1.08 (3H, d, J=5.9Hz), 1.20-1.58
(8H, m), 1.66-1.95 (5H, m), 2.10-2.60 (4H, m),
30 3.09-3.30 (1H, m), 3.58-4.60 (15H, m), 4.69-5.20
(10H, m), 5.24 (1H, d, J=4.5Hz), 5.51 (1H, d,
J=6.0Hz), 6.68-6.95 (4H, m), 7.04 (1H, d,
J=1.0Hz), 7.10-7.73 (7H, m), 7.73-7.90 (2H, m),
7.98 (1H, d, J=1.9Hz), 8.10 (1H, d, J=8.4Hz),
35 8.32 (1H, d, J=8.4Hz), 8.50 (1H, d, J=7.7Hz),

8.84 (1H, s)

FAB-MASS : $m/z = 1275$ ($M^+ + Na$)

Elemental Analysis Calcd. for $C_{55}H_{73}N_8O_{22}SNa \cdot 5H_2O$:

C 50.38, H 6.38, N 8.55

5

Found : C 49.98, H 6.37, N 8.41

Example 22

IR (KBr) : 3340, 2931, 1664, 1627, 1531, 1444, 1278,
1047 cm^{-1}

10 NMR (DMSO- d_6 , δ) : 0.86 (3H, t, $J=6.6Hz$), 0.96 (3H,
d, $J=6.8Hz$), 1.08 (3H, d, $J=5.9Hz$), 1.2-1.4 (6H,
m), 1.5-1.7 (2H, m), 1.7-2.1 (3H, m), 2.2-2.4
(3H, m), 2.6-2.7 (3H, m), 3.1-3.2 (1H, m), 3.7-
15 4.6 (13H, m), 4.78 (1H, d, $J=6.0Hz$), 4.8-5.1
(1H, m), 5.09 (1H, d, $J=5.6Hz$), 5.16 (1H, d,
 $J=3.2Hz$), 5.24 (1H, d, $J=4.4Hz$), 5.52 (1H, d,
 $J=6.0Hz$), 6.73 (1H, d, $J=8.2Hz$), 6.83 (2H, d,
 $J=8.3Hz$), 7.05 (1H, s), 7.3-7.5 (5H, m), 7.65
(2H, d, $J=8.2Hz$), 7.74 (2H, d, $J=8.4Hz$), 7.98
20 (2H, d, $J=8.4Hz$), 8.11 (1H, d, $J=8.4Hz$), 8.31
(1H, d, $J=8.4Hz$), 8.79 (1H, d, $J=7.7Hz$), 8.84
(1H, s)

FAB-MASS : $m/z = 1245$ ($M^+ + Na$)

Elemental Analysis Calcd. for $C_{54}H_{71}N_8O_{21}SNa \cdot 4H_2O$:

25

C 50.07, H 6.15, N 8.65

Found : C 50.26, H 6.44, N 8.67

Example 23

30 NMR (DMSO- d_6 , δ) : 0.91 (3H, t, $J=6.7Hz$), 0.96 (3H,
d, $J=6.8Hz$), 1.05 (3H, d, $J=5.6Hz$), 1.2-1.5 (6H,
m), 1.6-2.1 (5H, m), 2.1-2.7 (4H, m), 3.0-3.5
(9H, m), 3.6-4.5 (15H, m), 4.6-5.6 (11H, m),
6.73 (1H, d, $J=8.2Hz$), 6.8-6.9 (4H, m), 6.95
(2H, d, $J=8.6Hz$), 7.02 (2H, d, $J=9.2Hz$), 7.04
35 (1H, s), 7.2-7.5 (3H, m), 7.82 (2H, d, $J=8.6Hz$),

8.06 (1H, d, J=8Hz), 8.25 (1H, d, J=6.7Hz), 8.43
(1H, d, J=6.7Hz), 8.85 (1H, s)

IR (KBr) : 3350, 1668, 1629, 1510 cm^{-1}

FAB-MASS : $m/z = 1345$ (M+Na)

5 Elemental Analysis Calcd. for $\text{C}_{58}\text{H}_{79}\text{N}_{10}\text{O}_{22}\text{SNa}\cdot 6\text{H}_2\text{O}$:
C 48.67, H 6.41, N 9.78
Found : C 48.80, H 6.46, N 9.82

Example 24

10 Major product

IR (KBr) : 3350, 1668, 1631, 1047 cm^{-1}

NMR (DMSO-d_6 , δ) : 0.96 (3H, d, J=6.7Hz), 1.08 (3H,
d, J=5.7Hz), 1.2-1.6 (10H, m), 1.6-2.4 (8H, m),
2.5-2.7 (1H, m), 3.18 (1H, m), 3.21 (3H, s),
15 3.29 (2H, t, J=6.4Hz), 3.6-3.83 (2H, m), 3.83-
4.6 (13H, m), 4.7-5.4 (11H, m), 5.51 (1H, d,
J=5.9Hz), 6.73 (1H, d, J=8.2Hz), 6.83 (1H, d,
J=8.2Hz), 6.85 (1H, s), 7.04 (2H, d, J=8.4Hz),
7.06 (1H, s), 7.31 (1H, s), 7.2-7.5 (2H, m),
20 7.67 (2H, d, J=8.4Hz), 7.71 (2H, d, J=8.4Hz),
7.96 (2H, d, J=8.4Hz), 8.06 (1H, d, J=8Hz), 8.25
(1H, d, J=6.7Hz), 8.74 (1H, d, J=6.7Hz), 8.84
(1H, s)

FAB-MASS : $m/z = 1319$ (M+Na)

25 Elemental Analysis Calcd. for $\text{C}_{57}\text{H}_{77}\text{N}_8\text{O}_{23}\text{SNa}\cdot 4\text{H}_2\text{O}$:
C 49.99, H 6.26, N 8.18
Found : C 49.74, H 6.27, N 8.06

Minor product

30 IR (KBr) : 3350, 1668, 1631 cm^{-1}

NMR (DMSO-d_6 , δ) : 0.96 (3H, d, J=6.7Hz), 1.08 (3H,
d, J=5.7Hz), 1.2-1.6 (6H, m), 1.6-2.1 (7H, m),
2.1-2.5 (3H, m), 2.5-2.7 (1H, m), 3.18 (1H, m),
3.6-3.8 (2H, m), 3.8-4.6 (13H, m), 4.6-5.2 (12H,
35 m), 5.26 (1H, d, J=4.6Hz), 5.53 (1H, d,

J=5.8Hz), 5.6-6.0 (1H, m), 6.73 (1H, d,
J=8.2Hz), 6.83 (1H, d, J=8.3Hz), 6.85 (1H, s),
7.04 (2H, d, J=8.5Hz), 7.06 (1H, s), 7.30 (1H,
s), 7.2-7.5 (2H, m), 7.68 (2H, d, J=8.5Hz), 7.72
5 (2H, d, J=8.5Hz), 7.96 (2H, d, J=8.5Hz), 8.06
(1H, d, J=8Hz), 8.25 (1H, d, J=6.7Hz), 8.74 (1H,
d, J=6.7Hz), 8.85 (1H, s)

FAB-MASS : $m/z = 1287$ (M+Na)

Elemental Analysis Calcd. for $C_{56}H_{73}N_8NaO_{22}S \cdot 7H_2O$:

10 C 48.34, H 6.30, N 8.05
Found : C 48.19, H 6.19, N 7.99

Example 25

15 IR (KBr) : 3350, 2935, 2873, 1668, 1629, 1538, 1506,
1438, 1257, 1049 cm^{-1}

NMR (DMSO- d_6 , δ) : 0.9-1.0 (6H, m), 1.08 (3H, d,
J=5.7Hz), 1.2-1.6 (4H, m), 1.6-2.0 (5H, m), 2.1-
2.4 (3H, m), 2.5-2.6 (1H, m), 3.1-3.2 (1H, m),
3.6-4.6 (15H, m), 4.7-5.2 (10H, m), 5.26 (1H, d
20 J=4.5Hz), 5.55 (1H, d, J=5.9Hz), 6.7-6.9 (3H,
m), 7.0-7.6 (7H, m), 7.85 (2H, d, J=8.6Hz), 7.9-
8.2 (4H, m), 8.26 (1H, d, J=7.7Hz), 8.8-9.0 (2H,
m)

FAB-MASS : $m/z = 1314.3$ (M+Na)⁺

25 Elemental Analysis Calcd. for $C_{56}H_{70}N_9O_{23}NaS \cdot 7H_2O$:

C 47.42, H 5.97, N 8.89
Found : C 47.33, H 5.85, N 8.73

Example 26

30 To a solution of The Starting Compound (1 g) and
succinimido 4-(4-octyloxyphenyl)piperazine-1-carboxylate
(0.45 g) in N,N-dimethylformamide (10 ml) was added 4-
dimethylaminopyridine (0.141 g), and stirred for 5 days at
50°C. The reaction mixture was pulverized with ethyl
35 acetate. The precipitate was collected by filtration, and

dried under reduced pressure. The powder was dissolved in water, and subjected to column chromatography on ion exchange resin (DOWEX-50WX4) eluting with water. The fractions containing the object compound were combined, and subjected to column chromatography on ODS (YMC-gel-ODS-AM-S-50) eluting with 50% acetonitrile aqueous solution. The fractions containing the object compound were combined, and evaporated under reduced pressure to remove acetonitrile. The residue was lyophilized to give crude The Object Compound (23). The powder of crude The Object Compound (23) was purified by preparative HPLC utilizing a C₁₈ μ Bondapak resin (Waters Associates, Inc.) which was eluted with a solvent system comprised of (acetonitrile-pH 3 phosphate buffer = 40:60) at a flow rate of 80 ml/minute using a Shimadzu LC-8A pump. The column was monitored by a UV detector set at 240 nm. The fractions containing the object compound were combined, and evaporated under reduced pressure to remove acetonitrile. The residue was subjected to column chromatography on ion exchange resin (DOWEX-50WX4) eluting with water. The fractions containing the object compound were combined, and subjected to column chromatography on ODS (YMC-gel-ODS-AM-S-50) eluting with 50% acetonitrile aqueous solution. The fractions containing the object compound were combined, and evaporated under reduced pressure to remove acetonitrile. The residue was lyophilized to give The Object Compound (23) (60 mg).

IR (KBr) : 3347, 1629, 1511, 1245 cm⁻¹

NMR (DMSO-d₆, δ) : 0.86 (3H, t, J=6.7Hz), 0.95 (3H, d, J=6.8Hz), 1.06 (3H, d, J=5.9Hz), 1.2-1.5 (10H, m), 1.55-1.92 (5H, m), 2.0-2.65 (4H, m), 2.8-3.05 (5H, m), 3.2-4.47 (17H, m), 4.6-5.6 (12H, m), 6.6-7.0 (7H, m), 7.03 (1H, s), 7.2-7.5 (3H, m), 7.9-8.3 (3H, m), 8.84 (1H, s)

FAB-MASS : m/z = 1297 (M⁺+Na)

Elemental Analysis Calcd. for $C_{54}H_{79}N_{10}O_{22}SNa \cdot 6H_2O \cdot CH_3CN$:

C 47.22, H 6.65, N 10.82

Found : C 47.58, H 7.05, N 10.85

5 Example 27

To a suspension of 1-hydroxybenzotriazole (0.53 g) and 2-(4-octyloxyphenoxy)acetic acid (1 g) in dichlormethane (30 ml) was added 1-ethyl-3-(3'-dimethylaminopropyl)carbodiimide hydrochloride (WSCD·HCl) (0.886 g), and stirred for 3 hours at ambient temperature. The reaction mixture was added to water. The organic layer was taken, and dried over magnesium sulfate. The magnesium sulfate was filtered off, and the filtrate was evaporated under reduced pressure to give 1-[2-(4-octyloxyphenoxy)acetyl]benzotriazole 3-oxide (892 mg). To a solution of The Starting Compound (1.79 g) and 1-[2-(4-octyloxyphenoxy)acetyl]benzotriazole 3-oxide (892 mg) in N,N-dimethylformamide (18 ml) was added 4-(N,N-dimethylamino)pyridine (0.297 g), and stirred for 12 hours at ambient temperature. The reaction mixture was pulverized with ethyl acetate. The precipitate was collected by filtration, and dried under reduced pressure. The powder was added to water, and subjected to ion-exchange column chromatography on DOWEX-50WX4, and eluted with water. The fractions containing the object compound were combined, and subjected to column chromatograph on ODS (YMC-gel·ODS-AM·S-50), and eluted with 50% methanol aqueous solution. The fractions containing the object compound were combined, and evaporated under reduced pressure to remove methanol. The residue was lyophilized to give The Object Compound (24) (1.75 g).

IR. (KBr) : 3350, 1666, 1629, 1228 cm^{-1}

NMR (DMSO- d_6 , δ) : 0.86 (3H, t, J=6.9Hz), 0.95 (3H, d, J=6.7Hz), 1.04 (3H, d, J=5.7Hz), 1.15-1.5 (10H, m), 1.55-2.0 (5H, m), 2.05-2.5 (4H, m),

3.16 (1H, m), 3.72 (2H, m), 3.88 (3H, t,
J=6.3Hz), 4.41 (2H, s), 3.93-4.6 (11H, m),
4.69-5.25 (10H, m), 5.28 (1H, d, J=4.3Hz), 5.57
(1H, d, J=5.7Hz), 6.73 (1H, d, J=8.2Hz), 6.8-7.0
(5H, m), 7.04 (1H, s), 7.09 (1H, s), 7.3-7.4
(2H, m), 7.92-8.17 (2H, m), 8.29 (1H, d,
J=7.5Hz), 8.84 (1H, s)

FAB-MASS : $m/z = 1243$ ($M^+ + Na$)

Elemental Analysis Calcd. for $C_{51}H_{73}N_8O_{23}SNa \cdot 4H_2O$:

C 47.36, H 6.31, N 8.66

Found : C 47.22, H 6.44, N 8.37

The Object Compounds (28) to (31) were obtained
according to a similar manner to that of Example 27.

Example 28

IR (KBr) : 3350, 2933, 1664, 1628, 1446, 1205,
1045 cm^{-1}

NMR (DMSO- d_6 , δ) : 0.8-1.1 (9H, m), 1.2-2.0 (19H, m),
2.1-2.3 (3H, m), 3.6-3.8 (4H, m), 3.9-4.4 (13H, m),
4.6-5.0 (8H, m), 5.07 (1H, d, J=5.6Hz), 5.14 (1H,
d, J=3.2Hz), 5.23 (1H, d, J=4.3Hz), 5.46 (1H, d,
J=6.7Hz), 6.7-6.9 (3H, m), 7.04 (1H, s), 7.2-7.5
(6H, m), 7.8-8.0 (3H, m), 8.05 (1H, d, J=8.4Hz),
8.2-8.4 (2H, m), 8.83 (1H, s)

FAB-MASS : $m/z = 1360$ ($M^+ + Na$)

Elemental Analysis Calcd. for $C_{59}H_{80}N_9O_{23}SNa \cdot 6H_2O$:

C 48.99, H 6.41, N 8.72

Found : C 48.92, H 6.37, N 8.64

Example 29

IR (KBr) : 3350, 2927, 1668, 1627, 1535, 1515, 1452,
1440, 1286, 1045 cm^{-1}

NMR (DMSO- d_6 , δ) : 0.83 (3H, t, J=6.7Hz), 0.95 (3H,
d, J=6.7Hz), 1.07 (3H, d, J=5.9Hz), 1.2-1.4

(12H, m), 1.6-2.0 (5H, m), 2.1-2.4 (3H, m), 2.6
(1H, m), 2.82 (2H, t, J=7.4Hz), 3.1-3.2 (1H, m),
3.6-4.5 (13H, m), 4.7-5.2 (11H, m), 5.4-5.6 (1H,
m), 6.72 (1H, d, J=8.2Hz), 6.82 (2H, d,
5 J=8.1Hz), 7.03 (1H, s), 7.2-7.4 (3H, m), 7.47
(1H, d, J=8.5Hz), 7.69 (1H, d, J=8.5Hz), 8.1-8.2
(2H, m), 8.23 (1H, d, J=8.4Hz), 8.62 (1H, d,
J=7.8Hz), 8.83 (1H, s)

FAB-MASS : $m/z = 1251 (M^+ + Na)$

10 Elemental Analysis Calcd. for $C_{52}H_{73}N_{10}O_{21}SNa \cdot 5H_2O$:

C 47.34, H 6.34, N 10.61

Found : C 47.30, H 6.45, N 10.45

Example 30

15 NMR (DMSO- d_6 , δ) : 0.86 (3H, t, J=6.8Hz), 0.96 (3H,
t, J=6.7Hz), 1.05 (3H, t, J=5.8Hz), 1.2-1.5
(10H, m), 1.6-2.0 (5H, m), 2.2-2.4 (3H, m), 2.5-
2.6 (1H, m), 3.1-3.2 (1H, m), 3.7-4.5 (15H, m),
4.7-5.0 (8H, m), 5.10 (1H, d, J=5.6Hz), 5.17
20 (1H, d, J=3.1Hz), 5.26 (1H, d, J=4.5Hz), 5.52
(1H, d, J=5.8Hz) 6.73 (1H, d, J=8.2Hz), 6.8-7.0
(3H, m), 7.04 (1H, s), 7.2-7.4 (3H, m), 8.0-8.3
(3H, m), 8.68 (1H, d, J=2.3Hz), 8.7-8.8 (1H, m),
8.85 (1H, m)

25 FAB-MASS : $m/z = 1214 (M^+ + Na)$

Elemental Analysis Calcd. for $C_{49}H_{70}N_9O_{22}SNa \cdot 4H_2O$:

C 46.55, H 6.22, N 9.97

Found : C 46.29, H 6.18, N 9.71

30 Example 31

IR (Nujol) : 3342, 2210, 1668, 1623 cm^{-1}

NMR (DMSO- d_6 , δ) : 0.88 (3H, t, J=6.7Hz), 0.97 (3H,
d, J=6.7Hz), 1.08 (3H, d, J=6.7Hz), 1.20-1.60
(8H, m), 1.60-2.00 (5H, m), 2.05-2.50 (4H, m),
35 3.05-3.30 (1H, m), 3.60-4.60 (15H, m), 4.65-5.18

(10H, m), 5.24 (1H, d, $J=4.5\text{Hz}$), 5.58 (1H, d, $J=6.0\text{Hz}$), 6.68-7.10 (4H, m), 7.15-7.65 (5H, m), 7.80-8.30 (6H, m), 8.84 (1H, s), 9.18 (1H, d, $J=7.7\text{Hz}$)

5 FAB-MASS : $m/z = 1273.5$ ($M^+ + \text{Na}$)

Example 32

To a solution of 6-heptyloxy-2-naphthoic acid (0.358 g) and triethylamine (0.174 ml) in *N,N*-dimethylformamide (10 ml) was added diphenylphosphoryl azide (0.4 ml), and stirred for an hour at ambient temperature. Then, the reaction mixture was stirred for an hour at 100°C . After cooling, to the reaction mixture was added The Starting Compound (1 g) and 4-(*N,N*-dimethylamino)pyridine (0.140 g), and stirred for 10 hours at ambient temperature. The reaction mixture was pulverized with ethyl acetate. The precipitate was collected by filtration, and dried under reduced pressure. The powder was dissolved in water, and subjected to column chromatography on ion exchange resin (DOWEX-50WX4) eluting with water. The fractions containing the object compound were combined, and subjected to column chromatography on ODS (YMC-gel-ODS-AM-S-50) eluting with 50% acetonitrile aqueous solution. The fractions containing the object compound were combined, and evaporated under reduced pressure to remove acetonitrile. The residue was lyophilized to give The Object Compound (29) (0.832 g).

IR (KBr) : 3350, 1664, 1629, 1546, 1240 cm^{-1}

30 NMR ($\text{DMSO}-d_6$, δ) : 0.88 (3H, t, $J=6.6\text{Hz}$), 0.97 (3H, d, $J=6.7\text{Hz}$), 1.08 (3H, d, $J=5.9\text{Hz}$), 1.2-1.55 (8H, m), 1.55-2.0 (5H, m), 2.1-2.5 (4H, m), 3.18 (1H, m), 3.6-3.8 (3H, m), 3.9-4.5 (13H, m), 4.7-4.95 (3H, m), 5.0-5.3 (7H, m), 5.59 (1H, d, $J=5.8\text{Hz}$), 6.52 (1H, d, $J=8.1\text{Hz}$), 6.73 (1H, d, $J=8.2\text{Hz}$), 6.83 (1H, d, $J=8.2\text{Hz}$), 6.90 (1H, s),

7.0-7.15 (3H, m), 7.20 (1H, s), 7.27-7.4 (3H, m), 7.6-7.7 (2H, m), 7.87 (1H, s), 7.95-8.2 (2H, m), 8.69 (1H, s), 8.85 (1H, s)

FAB-MS : $m/z = 1264$ ($M^+ + Na$)

5 Elemental Analysis Calcd. for $C_{53}H_{72}N_9O_{22}SNa \cdot 5H_2O$:
C 47.78, H 6.20, N 9.46
Found : C 47.65, H 6.42, N 9.34

10 The Object Compound (33) was obtained according to a similar manner to that of Example 32.

Example 33

IR (KBr) : 3350, 1666, 1629, 1537, 1240 cm^{-1}

15 NMR (DMSO- d_6 , δ) : 0.87 (3H, t, $J=6.7Hz$), 0.97 (3H, d, $J=6.7Hz$), 1.09 (3H, d, $J=5.8Hz$), 1.2-1.55 (8H, m), 1.55-2.0 (5H, m), 2.07-2.6 (4H, m), 3.18 (1H, m), 3.6-3.85 (3H, m), 3.9-4.5 (13H, m), 4.7-4.98 (3H, m), 5.0-5.3 (7H, m), 5.57 (1H, d, $J=5.9Hz$), 6.50 (1H, d, $J=8.1Hz$), 6.73 (1H, d, $J=8.2Hz$), 6.82 (1H, dd, $J=8.2$ and $1.7Hz$), 6.87 (1H, s), 6.97 (2H, d, $J=8.8Hz$), 7.05 (1H, d, $J=1.7Hz$), 7.10 (1H, s), 7.23-7.43 (2H, m), 7.38 (2H, d, $J=8.8Hz$), 7.50 (2H, d, $J=8.8Hz$), 7.52 (2H, d, $J=8.8Hz$), 8.0-8.15 (2H, m), 8.65 (1H, s), 8.84 (1H, s)

25 FAB-MASS : $m/z = 1290$ ($M^+ + Na$)

Elemental Analysis Calcd. for $C_{55}H_{74}N_9O_{22}SNa \cdot 7H_2O$:
C 47.38, H 6.36, N 9.04
Found : C 47.67, H 6.53, N 9.03

30

Example 34

A solution of The Starting Compound (2.45 g), 3-[4-(4-pentylphenyl)phenyl]propionic acid (0.90 g), 1-ethyl-3-(3'-dimethylaminopropyl)carbodiimide hydrochloride (WSCD-HCl) (0.59 g) and triethylamine (0.43 ml) in N,N-

35

dimethylformamide (50 ml) was stirred for 15 hours at ambient temperature. The reaction mixture was diluted with ethyl acetate, and the resultant precipitate was collected by filtration, and washed in turn with ethyl acetate and diisopropyl ether, and dried under reduced pressure. The powder was dissolved in water, and was subjected to column chromatography on ion exchange resin (DOWEX-50WX4 (Na form, 50 ml)) eluting with water. The fractions containing the object compound were combined, and subjected to reversed phase chromatography on ODS (YMC-gel-ODS-AM-S-50, 50 ml) eluting with (water : acetonitrile = 10:0 - 7:3, V/V). The fractions containing the object compound were combined, and evaporated under reduced pressure to remove acetonitrile. The residue was lyophilized to give The Object Compound (31) (1.53 g).

IR (Nujol) : 3351, 2212, 1668, 1627 cm^{-1}

NMR (DMSO- d_6 , δ) : 0.87 (3H, t, $J=6.5\text{Hz}$), 0.96 (3H, d, $J=6.7\text{Hz}$), 1.08 (3H, d, $J=5.8\text{Hz}$), 1.20-1.50 (4H, m), 1.50-2.00 (5H, m), 2.03-2.55 (4H, m), 2.62 (2H, t, $J=7.5\text{Hz}$), 3.17 (1H, t, $J=8.4\text{Hz}$), 3.55-4.57 (15H, m), 4.65-5.13 (9H, m), 5.16 (1H, d, $J=3.2\text{Hz}$), 5.24 (1H, d, $J=4.5\text{Hz}$), 5.58 (1H, d, $J=5.8\text{Hz}$), 6.67-6.90 (3H, m), 6.93-7.10 (2H, m), 7.15-7.50 (4H, m), 7.50-7.90 (6H, m), 8.06 (1H, d, $J=8.4\text{Hz}$), 8.15 (1H, d, $J=7.7\text{Hz}$), 8.84 (1H, s), 9.19 (1H, d, $J=7.1\text{Hz}$)

FAB-MASS : $m/z = 1255$ ($\text{M}^+ + \text{Na}$)

Elemental Analysis Calcd. for $\text{C}_{55}\text{H}_{69}\text{N}_8\text{O}_{21}\text{SNa}\cdot 4\text{H}_2\text{O}$:

C 50.61, H 5.95, N 8.58

Found : C 50.47, H 6.00, N 8.54

Example 35

To a suspension of 1-hydroxybenzotriazole (501 mg) and 4-(4-heptylphenyl)benzoic acid (1 g) in dichloromethane (30 ml) was added 1-ethyl-3-(3'-

dimethylaminopropyl)carbodiimide hydrochloride (WSCD·HCl) (839 mg), and stirred for 3 hours at ambient temperature. The reaction mixture was added to water. The organic layer was separated, and dried over magnesium sulfate.

5 The magnesium sulfate was filtered off, and the filtrate was evaporated under reduced pressure to give 1-[4-(4-heptylphenyl)benzoyl]benzotriazole 3-oxide. To a solution of The Starting Compound (2.49 g) and 1-[4-(4-

10 dimethylformamide (25 ml) was added 4-(N,N-dimethylamino)pyridine (381 mg), and stirred for 12 hours at ambient temperature. The reaction mixture was pulverized with ethyl acetate. The precipitate was collected by filtration, and dried under reduced pressure.

15 The residue was dissolved in water, and subjected to column chromatography on ion exchange resin (DOWEX-50WX4) eluting with water. The fraction containing the object compound were combined, and subjected to column chromatography on ODS (YMC-gel-ODS-AM-S-50) eluting with
20 30% acetonitrile aqueous solution. The fractions containing the object compound were combined, and evaporated under reduced pressure to remove acetonitrile. The residue was lyophilized to give The Object Compound (32) (1.99 g).

25 IR (Nujol) : 3350, 2852, 1749, 1621, 1457, 1376, 1045 cm^{-1}

NMR (DMSO- d_6 , δ) : 0.86 (3H, t, $J=6.7\text{Hz}$), 0.96 (3H, d, $J=6.7\text{Hz}$), 1.08 (3H, d, $J=5.9\text{Hz}$), 1.5-1.7 (2H, m), 1.7-2.2 (3H, m), 2.2-2.5 (3H, m), 2.6-2.8
30 (3H, m), 3.1-3.2 (1H, m), 3.7-4.6 (13H, m), 4.7-5.2 (8H, m), 5.12 (1H, d, $J=5.5\text{Hz}$), 5.18 (1H, d, $J=2.9\text{Hz}$), 5.27 (1H, d, $J=4.4\text{Hz}$), 5.54 (1H, d, $J=5.8\text{Hz}$), 6.7-6.9 (3H, m), 7.05 (1H, s), 7.2-7.4 (5H, m), 7.65 (2H, d, $J=8.0\text{Hz}$), 7.74
35 (2H, d, $J=8.3\text{Hz}$), 7.98 (2H, d, $J=8.3\text{Hz}$), 8.11

(1H, d, J=8.7Hz), 8.28 (1H, d, J=8.4Hz), 8.78
(1H, d, J=7.3Hz), 8.85 (1H, s)

FAB-MASS : m/z = 1259 (M⁺+Na)

Elemental Analysis Calcd. for C₅₅H₇₃N₈O₂₁SNa·5H₂O :

C 49.77, H 6.30, N 8.44

Found : C 49.98, H 6.44, N 8.41

The Object Compounds (36) to (107) were obtained
according to a similar manner to that of Example 1.

Example 36

IR (KBr) : 3350, 1675.8, 1629.6, 1515.8 cm⁻¹

NMR (DMSO-d₆, δ) : 0.86 (6H, d, J=6.6Hz), 0.96 (3H, d,
J=6.6Hz), 1.06 (3H, d, J=5.7Hz), 1.1-1.3 (2H, m),
1.4-2.0 (6H, m), 2.0-2.7 (4H, m), 3.1-3.5 (9H, m),
3.66 (2H, t, J=7.3Hz), 3.6-4.5 (13H, m), 4.7-5.6
(12H, m), 6.73 (1H, d, J=8.3Hz), 6.82 (1H, d,
J=8.3Hz), 6.8-6.9 (1H, m), 7.02 (2H, d, J=9.0Hz),
7.04 (1H, s), 7.11 (2H, d, J=9.0Hz), 7.2-7.6 (3H,
m), 7.50 (2H, d, J=9.0Hz), 7.82 (2H, d, J=9.0Hz),
8.1 (1H, d, J=8.5Hz), 8.28 (1H, d, J=8.5Hz), 8.33
(1H, s), 8.45 (1H, d, J=7.0Hz), 8.84 (1H, s)

FAB-MASS : m/z = 1412 (M+Na)

Elemental Analysis Calcd. for C₆₀H₈₀N₁₃O₂₂SNa·9H₂O :

C 46.42, H 6.36, N 11.73

Found : C 46.64, H 6.43, N 11.62

Example 37

IR (KBr) : 3350, 1668.1, 1629.6, 1268.9 cm⁻¹

NMR (DMSO-d₆, δ) : 0.85 (3H, t, J=6.6Hz), 0.96 (3H, d,
J=6.7Hz), 1.07 (3H, d, J=5.9Hz), 1.2-1.4 (10H, m),
1.4-2.0 (5H, m), 2.0-2.5 (4H, m), 2.61 (2H, t,
J=7.2Hz), 3.1-3.3 (1H, m), 3.6-4.5 (13H, m), 4.40
(2H, s), 4.6-5.3 (11H, m), 5.60 (1H, d, J=5.8Hz),
6.73 (1H, d, J=8.2Hz), 6.82 (1H, d, J=8.2Hz), 6.6-

6.9 (1H, m), 7.04 (1H, s), 7.0-7.1 (1H, m), 7.32
(2H, d, J=8.5Hz), 7.2-7.5 (2H, m), 7.58 (2H, d,
J=8.5Hz), 7.93 (1H, d, J=7Hz), 8.04 (1H, d,
J=9.4Hz), 8.41 (1H, s), 8.44 (1H, d, J=9.4Hz), 8.84
(1H, s)

FAB-MASS : m/z = 1294 (M+Na)

Elemental Analysis Calcd. for $C_{53}H_{74}N_{11}O_{22}SNa \cdot 7H_2O$:

C 45.52, H 6.34, N 11.02

Found : C 45.47, H 6.27, N 10.93

Example 38

Major product

IR (KBr) : 3349.7, 1670.1, 1627.6, 1508.1 cm^{-1}

NMR (DMSO- d_6 , δ) : 0.96 (3H, d, J=6.6Hz), 1.06 (3H, d,
J=5.7Hz), 1.2-1.6 (8H, m), 1.6-2.1 (5H, m), 2.1-2.7
(4H, m), 3.0-3.2 (5H, m), 3.21 (3H, s), 3.30 (2H,
t, J=6.5Hz), 3.3-3.5 (4H, m), 3.6-4.5 (15H, m),
4.7-5.3 (11H, m), 5.49 (1H, d, J=5.9Hz), 6.73 (1H,
d, J=8.3Hz), 6.8-6.9 (4H, m), 6.95 (2H, d,
J=9.2Hz), 7.01 (2H, d, J=8.5Hz), 7.04 (1H, s), 7.20
(1H, s), 7.2-7.5 (2H, m), 7.81 (2H, d, J=8.5Hz),
8.09 (1H, d, J=8.7Hz), 8.28 (1H, d, J=8.7Hz), 8.45
(1H, d, J=6.7Hz), 8.84 (1H, s)

FAB-MASS : m/z = 1389 (M+Na)

Elemental Analysis Calcd. for $C_{60}H_{83}N_{10}O_{23}SNa \cdot 8H_2O$:

C 47.68, H 6.60, N 9.27

Found : C 47.83, H 6.72, N 9.27

Minor product

IR (KBr) : 3338.2, 1646.9, 1511.9 cm^{-1}

NMR (DMSO- d_6 , δ) : 0.96 (3H, d, J=6.7Hz), 1.06 (3H, d,
J=5.7Hz), 1.3-1.6 (4H, m), 1.6-2.7 (11H, m), 3.0-
3.2 (5H, m), 3.3-3.5 (4H, m), 3.6-4.5 (15H, m),
4.7-5.3 (13H, m), 5.48 (1H, d, J=5.9Hz), 5.7-6.0
(1H, m), 6.73 (1H, d, J=8.2Hz), 6.8-6.9 (4H, m),

6.94 (2H, d, J=9.3Hz), 7.01 (2H, d, J=8.6Hz), 7.04
(1H, s), 7.2-7.5 (3H, m), 7.81 (2H, d, J=8.6Hz),
8.06 (1H, d, J=8.7Hz), 8.25 (1H, d, J=8.7Hz), 8.42
(1H, d, J=6.7Hz), 8.84 (1H, s)

5 FAB-MASS : m/z = 1357 (M+Na)

Elemental Analysis Calcd. for $C_{59}H_{79}N_{10}O_{22}SNa \cdot 9H_2O$:

C 47.32, H 6.53, N 9.35

Found : C 47.08, H 6.66, N 9.25

10 Example 39

IR (KBr) : 3350, 1670.1, 1631.5, 1510.0, 1234.2 cm^{-1}

NMR (DMSO- d_6 , δ) : 0.87 (3H, t, J=6.7Hz), 0.96 (3H, d,
J=6.7Hz), 1.06 (3H, d, J=5.6Hz), 1.2-1.5 (8H, m),
1.6-2.1 (5H, m), 2.1-2.7 (4H, m), 3.0-3.3 (5H, m),
15 3.3-3.5 (4H, m), 3.6-3.8 (2H, m), 3.88 (2H, d,
J=6.4Hz), 3.8-4.5 (11H, m), 4.7-5.1 (8H, m), 5.10
(1H, d, J=5.6Hz), 5.16 (1H, d, J=3.1Hz), 5.25 (1H,
d, J=4.5Hz), 5.48 (1H, d, J=5.9Hz), 6.73 (1H, d,
J=8.2Hz), 6.8-6.9 (4H, m), 6.94 (2H, d, J=9.3Hz),
20 7.01 (2H, d, J=8.7Hz), 7.04 (1H, s), 7.2-7.5 (3H,
m), 7.81 (2H, d, J=8.7Hz), 8.06 (1H, d, J=8Hz),
8.25 (1H, d, J=6.7Hz), 8.43 (1H, d, J=6.7Hz), 8.85
(1H, s)

FAB-MASS : m/z = 1359 (M+Na)

25 Elemental Analysis Calcd. for $C_{59}H_{81}N_{10}O_{22}SNa \cdot 5H_2O$:

C 49.64, H 6.43, N 9.81

Found : C 49.49, H 6.54, N 9.72

Example 40

30 IR (KBr) : 3355.5, 1670.1, 1627.6, 1510.0 1236.1 cm^{-1}

NMR (DMSO- d_6 , δ) : 0.89 (6H, d, J=6.5Hz), 0.96 (3H, d,
J=6.7Hz), 1.05 (3H, d, J=5.7Hz), 1.2-1.4 (2H, m),
1.5-2.1 (6H, m), 2.1-2.7 (4H, m), 3.0-3.6 (9H, m),
3.6-4.5 (15H, m), 4.5-5.4 (12H, m), 6.73 (1H, d,
35 J=8.2Hz), 6.8-6.9 (4H, m), 6.96 (2H, d, J=9.6Hz),

7.02 (2H, d, J=8.7Hz), 7.05 (1H, s), 7.2-7.5 (3H, m), 7.82 (2H, d, J=8.7Hz), 8.08 (1H, d, J=8Hz), 8.27 (1H, d, J=6.7Hz), 8.46 (1H, d, J=6.7Hz), 8.85 (1H, s)

5 FAB-MASS : m/z = 1345 (M+Na)

Elemental Analysis Calcd. for $C_{58}H_{79}N_{10}O_{22}SNa \cdot 8H_2O$:

C 47.47, H 6.52, N 9.54

Found : C 47.47, H 6.54, N 9.51

10 Example 41

IR (KBr) : 3347.8, 1668.1, 1629.6, 1510.0, 1234.2 cm^{-1}

15 NMR (DMSO- d_6 , δ) : 0.89 (3H, t, J=7.0Hz), 0.96 (3H, d, J=6.7Hz), 1.05 (3H, d, J=5.8Hz), 1.2-1.5 (4H, m), 1.6-2.1 (5H, m), 2.1-2.7 (4H, m), 3.0-3.6 (9H, m), 3.6-3.8 (2H, m), 3.8-4.5 (13H, m), 4.7-5.6 (12H, m), 6.73 (1H, d, J=8.2Hz), 6.8-6.9 (4H, m), 6.96 (2H, d, J=8.7Hz), 7.02 (2H, d, J=9.0Hz), 7.04 (1H, s), 7.2-7.5 (3H, m), 7.82 (2H, d, J=8.7Hz), 8.07 (1H, d, J=8Hz), 8.27 (1H, d, J=6.7Hz), 8.45 (1H, d, J=6.7Hz), 8.85 (1H, s)

20 FAB-MASS : m/z = 1331 (M+Na)

Elemental Analysis Calcd. for $C_{57}H_{77}N_{10}O_{22}SNa \cdot 6H_2O$:

C 48.30, H 6.33, N 9.88

Found : C 48.20, H 6.58, N 10.03

25 Example 42

Mixture product

IR (KBr) : 3344, 1670.1, 1631.5 cm^{-1}

30 NMR (DMSO- d_6 , δ) : 0.96 (3H, d, J=6.7Hz), 1.08 (3H, d, J=5.9Hz), 1.2-1.5 (8H, m), 1.6-2.1 (7H, m), 2.1-2.7 (4H, m), 3.1-3.3 (1H, m), 3.6-4.5 (15H, m), 4.45 and 4.70 (2H, t, J=7.1Hz), 4.6-5.3 (11H, m), 5.52 (1H, d, J=5.9Hz), 6.73 (1H, d, J=8.2Hz), 6.83 (1H, d, J=8.2Hz), 6.85 (1H, s), 7.03 (2H, d, J=8.6Hz), 7.05 (1H, s), 7.2-7.5 (3H, m), 7.68 (2H, d,

35

J=8.6Hz), 7.71 (2H, d, J=8.4Hz), 7.96 (2H, d,
J=8.4Hz), 8.12 (1H, d, J=8.5Hz), 8.30 (1H, d,
J=7.0Hz)

FAB-MASS : m/z = 1357 (M+Na)

5 Elemental Analysis Calcd. for $C_{57}H_{75}N_{12}O_{22}SNa \cdot 4H_2O$:
C 48.64, H 5.94, N 11.94
Found : C 48.91, H 5.88, N 11.86

Example 43

10 IR (KBr) : 3350, 1666.2, 1651.5 cm^{-1}
NMR (DMSO- d_6 , δ) : 0.96 (3H, d, J=6.7Hz), 1.05 (6H, d,
J=6.3Hz), 1.06 (3H, d, J=5.7Hz), 1.2-1.6 (10H, m),
1.6-2.1 (7H, m), 2.1-2.7 (6H, m), 2.8-3.0 (2H, m),
3.0-3.2 (1H, m), 3.4-3.7 (2H, m), 3.6-3.8 (2H, m),
15 3.8-4.5 (13H, m), 4.7-5.6 (12H, m), 6.73 (1H, d,
J=8.2Hz), 6.8-7.0 (2H, m), 7.03 (2H, d, J=8.7Hz),
7.06 (1H, s), 7.2-7.5 (3H, m), 7.67 (2H, d,
J=8.7Hz), 7.71 (2H, d, J=8.4Hz), 7.96 (2H, d,
J=8.4Hz), 8.04 (1H, d, J=8.5Hz), 8.31 (1H, d,
20 J=8.5Hz), 8.73 (1H, d, J=7.0Hz), 8.90 (1H, s)
FAB-MASS : m/z = 1402 (M+Na)

Example 44

25 IR (KBr pelet) : 3350, 2929, 2856, 1670, 1631, 1510,
1243, 1045 cm^{-1}
NMR (DMSO- d_6 , δ) : 0.86 (3H, t, J=6.8Hz), 0.96 (3H, d,
J=6.7Hz), 1.06 (3H, d, J=5.7Hz), 1.6-2.0 (5H, m),
2.2-2.5 (5H, m), 2.6-2.7 (1H, m), 3.0-3.3 (5H, m),
3.6-4.5 (19H, m), 4.77 (2H, d, J=5.9Hz), 4.8-5.1
30 (6H, m), 5.10 (1H, d, J=5.6Hz), 5.17 (1H, d,
J=3.1Hz), 5.25 (1H, d, J=4.5Hz), 5.50 (1H, d,
J=5.8Hz), 6.7-7.0 (8H, m), 7.04 (1H, s), 7.2-7.4
(3H, m), 8.0-8.2 (2H, m), 8.26 (1H, d, J=8.0Hz),
8.55 (1H, d, J=7.3Hz), 8.67 (1H, d, J=1.2Hz), 8.85
35 (1H, s)

FAB-MASS : $m/z = 1374.3$ ($M+Na^+$)

Elemental Analysis Calcd. for $C_{59}H_{82}N_{11}O_{22}NaS \cdot 5.5H_2O$:

C 48.82, H 6.46, N 10.61

Found : C 48.89, H 6.74, N 10.50

5

Example 45

IR (KBr) : 3350, 2935, 1668, 1623, 1538, 1257, 1174,
1047 cm^{-1}

10 NMR (DMSO- d_6 , δ) : 0.8-1.1 (6H, m), 1.09 (3H, d,
J=5.7Hz), 1.2-1.6 (6H, m), 1.7-2.1 (5H, m), 2.2-2.4
(3H, m), 2.5-2.6 (1H, m), 3.6-3.8 (2H, m), 3.8-4.6
(14H, m), 4.8-5.2 (7H, m), 5.18 (1H, d, J=3.1Hz),
5.26 (1H, d, J=4.5Hz), 5.54 (1H, d, J=5.8Hz), 6.7-
15 7.5 (9H, m), 7.82 (1H, d, J=8.5Hz), 7.96 (1H, d,
J=8.7Hz), 8.1-8.4 (5H, m), 8.8-9.0 (2H, m)

FAB-MASS : $m/z = 1302.6$ ($M+Na^+$)

Elemental Analysis Calcd. for $C_{55}H_{70}N_9O_{23}SNa \cdot 6H_2O$:

C 47.58, H 5.95, N 9.08

Found : C 47.46, H 6.04, N 9.05

20

Example 46

IR (KBr) : 3355, 2958, 1670, 1627, 1521, 1247,
1047 cm^{-1}

25 NMR (DMSO- d_6 , δ) : 0.9-1.0 (6H, m), 1.08 (3H, d,
J=5.6Hz), 1.4-1.6 (2H, m), 1.7-2.1 (5H, m), 2.1-2.4
(3H, m), 2.5-2.6 (1H, m), 3.1-3.3 (1H, m), 3.7-3.8
(2H, m), 3.9-4.6 (13H, m), 4.8-5.1 (8H, m), 5.11
(1H, d, J=5.6Hz), 5.17 (1H, d, J=3.1Hz), 5.26 (1H,
d, J=4.5Hz), 5.54 (1H, d, J=5.9Hz), 6.7-6.9 (3H,
30 m), 7.0-7.2 (3H, m), 7.3-7.5 (3H, m), 7.7-7.9 (8H,
m), 8.02 (2H, d, J=8.4Hz), 8.08 (1H, d, J=8.4Hz),
8.32 (1H, d, J=7.7Hz), 8.81 (1H, d, J=7.0Hz), 8.85
(1H, s)

FAB-MASS : $m/z = 1309.3$ ($M+Na$)⁺

35

Elemental Analysis Calcd. for $C_{58}H_{71}N_8O_{22}NaS \cdot 6H_2O$:

C 49.92, H 6.00, N 8.03

Found : C 49.92, H 5.97, N 8.03

Example 47

5 IR (KBr) : 3350, 2933, 1668, 1629, 1517, 1249,
1045 cm^{-1}

NMR (DMSO- d_6 , δ) : 0.88 (3H, t, $J=6.7\text{Hz}$), 0.96 (3H, d,
 $J=6.7\text{Hz}$), 1.08 (3H, d, $J=5.8\text{Hz}$), 1.7-2.7 (8H, m),
3.1-3.3 (1H, m), 3.6-4.5 (16H, m), 4.7-5.2 (8H, m),
10 5.18 (1H, d, $J=3.1\text{Hz}$), 5.27 (1H, d, $J=4.5\text{Hz}$), 5.56
(1H, d, $J=5.8\text{Hz}$), 6.7-7.0 (3H, m), 7.0-7.2 (3H, m),
7.2-7.5 (3H, m), 8.0-8.4 (6H, m), 8.85 (1H, s),
8.96 (1H, d, $J=7.0\text{Hz}$), 9.07 (1H, s)

FAB-MASS : $m/z = 1276.6$ ($M+\text{Na}^+$)

15 Elemental Analysis Calcd. for $\text{C}_{54}\text{H}_{72}\text{N}_9\text{O}_{22}\text{NaS}\cdot 5\text{H}_2\text{O}$:

C 48.25, H 6.15, N 9.38

Found : C 48.10, H 6.14, N 9.30

Example 48

IR (KBr) : 3350, 2931, 1668, 1629, 1537, 1049 cm^{-1}

20 NMR (DMSO- d_6 , δ) : 0.86 (3H, t, $J=6.9\text{Hz}$), 0.9-1.5
(16H, m), 1.6-2.4 (8H, m), 2.5-2.7 (1H, m), 3.1-3.3
(1H, m), 3.5-5.6 (25H, m), 6.6-7.4 (8H, m), 7.8-8.4
(6H, m), 8.7-9.0 (2H, m), 9.00 (1H, d, $J=2.4\text{Hz}$)

FAB-MASS : $m/z = 1331.4$ ($M+\text{Na}^+$)

25 Elemental Analysis Calcd. for $\text{C}_{56}\text{H}_{73}\text{N}_{10}\text{O}_{23}\text{NaS}\cdot 8\text{H}_2\text{O}$:

C 46.28, H 6.17, N 9.64

Found : C 46.50, H 6.27, N 9.65

Example 49

30 IR (KBr pelet) : 3300, 2931, 1668, 1650, 1629, 1538,
1515, 1268, 1049 cm^{-1}

NMR (DMSO- d_6 , δ) : 0.87 (3H, t, $J=6.6\text{Hz}$), 0.97 (3H, d,
 $J=6.7\text{Hz}$), 1.10 (3H, d, $J=5.6\text{Hz}$), 1.2-1.4 (6H, m),
1.5-1.7 (2H, m), 1.7-2.1 (3H, m), 2.1-2.4 (3H, m),
35 2.6-2.7 (3H, m), 3.1-3.2 (1H, m), 3.7-3.9 (2H, m),

3.9-4.5 (12H, m), 4.8-5.1 (7H, m), 5.11 (1H, d, J=5.5Hz), 5.18 (1H, d, J=3.1Hz), 5.27 (1H, d, J=4.5Hz), 5.55 (1H, d, J=5.8Hz), 6.7-7.0 (3H, m), 7.06 (1H, s), 7.3-7.5 (5H, m), 7.72 (2H, d, J=8.2Hz), 7.9-8.2 (5H, m), 8.3-8.4 (4H, m), 8.9-9.0 (2H, m)

FAB-MASS : $m/z = 1260.5$ ($M+Na^+$)

Elemental Analysis Calcd. for $C_{61}H_{74}N_9O_{22}SNa \cdot 6H_2O$:

C 50.58, H 5.98, N 8.70

Found : C 50.34, H 6.16, N 8.55

Example 50

IR (KBr) : 3369, 2958, 2935, 1670, 1629, 1525, 1473, 1247, 1047 cm^{-1}

NMR (DMSO- d_6 , δ) : 0.95 (3H, t, J=7.3Hz), 0.97 (3H, d, J=6.7Hz), 1.09 (3H, d, J=5.7Hz), 1.3-1.6 (2H, m), 1.7-2.1 (5H, m), 2.1-2.4 (3H, m), 2.5-2.6 (1H, m), 3.1-3.3 (1H, m), 3.7-4.6 (15H, m), 4.7-5.1 (8H, m), 5.10 (1H, d, J=5.6Hz), 5.18 (1H, d, J=3.1Hz), 5.26 (1H, d, J=4.4Hz), 5.56 (1H, d, J=5.7Hz), 6.7-7.0 (3H, m), 7.1-7.2 (3H, m), 7.2-7.4 (3H, m), 7.70 (2H, d, J=8.6Hz), 7.78 (2H, d, J=8.4Hz), 8.1-8.4 (6H, m), 8.85 (1H, s), 8.99 (1H, d, J=7.0Hz), 9.13 (1H, d, J=1.6Hz)

FAB-MASS : $m/z = 1310.1$ ($M+Na$)⁺

Elemental Analysis Calcd. for $C_{57}H_{70}N_9O_{22}NaS \cdot 7H_2O$:

C 47.20, H 6.12, N 8.69

Found : C 47.42, H 6.19, N 8.92

Example 51

IR (KBr) : 3351, 2937, 2875, 1670, 1627, 1533, 1245, 1047 cm^{-1}

NMR (DMSO- d_6 , δ) : 0.96 (3H, d, J=6.7Hz), 1.08 (3H, d, J=5.7Hz), 1.5-1.7 (2H, m), 1.7-2.1 (7H, m), 2.1-2.4 (3H, m), 2.5-2.6 (1H, m), 3.1-3.2 (1H, m), 3.7-

3.8 (2H, m), 3.9-4.6 (15H, m), 4.7-4.9 (3H, m),
5.0-5.1 (5H, m), 5.10 (1H, d, J=5.6Hz), 5.17 (1H,
d, J=3.1Hz), 5.26 (1H, d, J=4.5Hz), 5.52 (1H, d,
J=5.9Hz), 6.7-7.1 (9H, m), 7.2-7.5 (5H, m), 7.68
5 (2H, d, J=8.2Hz), 7.72 (2H, d, J=6.7Hz), 7.96 (2H,
d, J=8.2Hz), 8.06 (1H, d, J=8.4Hz), 8.28 (1H, d,
J=7.7Hz), 8.76 (1H, d, J=7.0Hz), 8.85 (1H, s)

FAB-MASS : $m/z = 1339.5$ ($M+Na^+$)

Elemental Analysis Calcd. for $C_{59}H_{73}N_8O_{23}NaS \cdot 7H_2O$:

C 49.09, H 6.08, N 7.76

Found : C 49.04, H 6.08, N 7.82

Example 52

IR (KBr) : 3350, 2954, 2937, 1670, 1631, 1440, 1257,
15 1047 cm^{-1}

NMR (DMSO- d_6 , δ) : 0.89 (3H, t, J=6.8Hz), 0.97 (3H, d,
J=6.7Hz), 1.09 (2H, d, J=5.8Hz), 1.2-1.5 (6H, m),
1.7-2.1 (5H, m), 2.1-2.4 (3H, m), 2.5-2.6 (1H, m),
3.1-3.2 (1H, m), 3.7-4.6 (15H, m), 4.7-5.3 (11H,
20 m), 5.5-5.6 (1H, m), 6.7-6.9 (1H, m), 7.0-7.5 (6H,
m), 8.0-8.4 (8H, m), 8.85 (1H, s), 8.96 (1H, d,
J=7.0Hz)

APCI-MASS : $m/z = 1329.0$ ($M+Na$)⁺

Elemental Analysis Calcd. for $C_{56}H_{71}N_{10}O_{23}NaS \cdot 6H_2O$:

C 47.52, H 5.91, N 9.90

Found : C 47.42, H 6.05, N 9.90

Example 53

IR (KBr) : 3350, 2952, 1666, 1629, 1537, 1519,
30 1255 cm^{-1}

NMR (DMSO- d_6 , δ) : 0.89 (3H, t, J=6.7Hz), 0.96 (3H, d,
J=6.4Hz), 1.08 (3H, d, J=5.6Hz), 1.7-2.4 (8H, m),
2.5-2.6 (1H, m), 3.7-4.5 (15H, m), 4.7-5.1 (8H, m),
5.11 (1H, d, J=5.5Hz), 5.17 (1H, d, J=3.1Hz), 5.26
35 (1H, d, J=3.1Hz), 5.56 (1H, d, J=5.7Hz), 6.73 (1H,

d, J=8.2Hz), 6.7-7.0 (2H, m), 7.05 (1H, s), 7.13
(2H, d, J=8.7Hz), 7.2-7.5 (3H, m), 7.97 (2H, d,
J=8.7Hz), 8.1-8.4 (6H, m), 8.85 (1H, s), 8.92 (1H,
d, J=7.0Hz)

5 FAB-MASS : $m/z = 1345.3$ (M+Na)⁺

Elemental Analysis Calcd. for

$C_{56}H_{71}N_{10}O_{22}S_2Na \cdot 8H_2O$:

C 45.84, H 5.98, N 9.55

Found : C 45.87, H 6.07, N 9.55

10

Example 54

IR (KBr pelet) : 3350, 2931, 1670, 1652, 1628, 1442,
1247, 1047 cm^{-1}

15 NMR (DMSO- d_6 , δ) : 0.86 (3H, t, J=6.6Hz), 0.97 (3H, d,
J=6.8Hz), 1.12 (3H, d, J=6.8Hz), 1.2-1.5 (10H, m),
1.7-2.0 (5H, m), 2.2-2.4 (3H, m), 2.5-2.6 (1H, m),
3.1-3.2 (1H, m), 3.72 (2H, br), 3.8-4.5 (17H, m),
4.7-5.2 (9H, m), 5.26 (1H, d, J=4.6Hz), 5.57 (1H,
d, J=5.7Hz), 6.7-7.1 (7H, m), 7.3-7.5 (3H, m), 7.66
20 (2H, d, J=8.7Hz), 8.10 (1H, d, J=7.6Hz), 8.17 (1H,
d, J=7.6Hz), 8.76 (1H, d, J=7.0Hz), 8.85 (1H, s)

FAB-MASS : $m/z = 1293$ (M+Na⁺)

Elemental Analysis Calcd. for $C_{54}H_{75}N_{10}O_{22}NaS \cdot 7H_2O$:

C 46.41, H 6.42, N 10.02

25 Found : C 46.51, H 6.43, N 9.95

Example 55

IR (KBr) : 3345, 2937, 1650, 1511, 1249, 1047 cm^{-1}

30 NMR (DMSO- d_6 , δ) : 0.91 (3H, t, J=7.0Hz), 0.96 (3H, t,
J=7.8Hz), 1.09 (3H, d, J=6.8Hz), 1.3-1.5 (4H, m),
1.6-2.1 (5H, m), 2.1-2.5 (3H, m), 2.5-2.6 (1H, m),
3.1-3.3 (1H, m), 3.7-3.9 (2H, m), 3.9-4.6 (13H, m),
4.79 (2H, d, J=5.9Hz), 4.8-4.9 (1H, m), 4.9-5.2
(5H, m), 5.10 (1H, d, J=5.9Hz), 5.17 (1H, d,
35 J=3.1Hz), 5.25 (1H, d, J=4.6Hz), 5.53 (1H, d,

J=5.9Hz), 6.7-7.0 (3H, m), 7.0-7.2 (3H, m), 7.19
(1H, s), 7.3-7.5 (3H, m), 7.7-8.1 (6H, m), 8.08
(1H, d, J=10.0Hz), 8.26 (1H, d, J=8.8Hz), 8.77 (1H,
m), 8.85 (1H, s), 13.32 (1H, s)

5 FAB-MASS : $m/z = 1314.0 (M+Na)^+$

Elemental Analysis Calcd. for $C_{56}H_{71}N_{10}O_{22}SNa \cdot 8H_2O$:

C 46.86, H 6.11, N 9.76

Found : C 46.93, H 5.87, N 9.74

10 Example 56

IR (KBr) : 3350, 2958, 2935, 2873, 1666, 1629, 1247,
1045 cm^{-1}

NMR (DMSO- d_6 , δ) : 0.9-1.1 (6H, m), 1.08 (3H, d,
J=6.0Hz), 1.4-1.6 (2H, m), 1.6-2.1 (5H, m), 2.1-2.4
15 (3H, m), 2.5-2.6 (1H, m), 3.1-3.3 (1H, m), 3.6-4.5
(15H, m), 4.7-5.1 (8H, m), 5.10 (1H, d, J=5.5Hz),
5.17 (1H, d, J=2.9Hz), 5.25 (1H, d, J=4.5Hz), 5.55
(1H, d, J=5.7Hz), 6.7-6.9 (3H, m), 7.0-7.5 (8H, m),
7.68 (2H, d, J=8.9Hz), 7.73 (2H, d, J=8.3Hz), 8.01
20 (2H, d, J=8.3Hz), 8.10 (1H, d, J=8.4Hz), 8.26 (1H,
d, J=7.7Hz), 8.8-9.0 (2H, m)

FAB-MASS : $m/z = 1299.5 (M+Na)^+$

Elemental Analysis Calcd. for $C_{56}H_{69}N_8O_{23}NaS \cdot 6H_2O$:

C 48.55, H 5.89, N 8.09

25 Found : C 48.52, H 5.94, N 8.07

Example 57

IR (KBr) : 3355.5, 1662.3, 1629.6, 1267.0 cm^{-1}

NMR (DMSO- d_6 , δ) : 0.88 (3H, t, J=6.8Hz), 0.93 (3H, d,
30 J=8.4Hz), 0.97 (3H, d, J=6.7Hz), 1.2-1.5 (4H, m),
1.5-1.95 (5H, m), 2.1-2.45 (4H, m), 2.5-2.7 (4H,
m), 3.17 (1H, m), 3.55-4.45 (14H, m), 4.6-5.3 (13H,
m), 5.56 (1H, d, J=5.6Hz), 6.72 (1H, d, J=8.1Hz),
6.75 (1H, s), 6.77 (1H, d, J=8.1Hz), 7.04 (1H, s),
35 7.10 (1H, s), 7.2-7.45 (10H, m), 7.53 (4H, d,

J=6.6Hz), 7.85 (1H, d, J=7Hz), 7.92 (1H, d, J=7Hz),
8.05 (1H, d, J=7Hz), 8.22 (1H, d, J=7Hz), 8.84 (1H,
s)

FAB-MASS : m/z = 1408 (M+Na)

5

Example 58

IR (KBr) : 3347.8, 1664.3, 1631.5, 1245.8 cm^{-1}

10 NMR (DMSO- d_6 , δ) : 0.86 (3H, t, J=6.6Hz), 0.96 (3H, d,
J=6.6Hz), 1.04 (3H, d, J=5.7Hz), 1.15-2.6 (21H,
m), 3.16 (1H, m), 3.5-4.5 (16H, m), 4.6-5.4 (13H,
m), 5.47 (1H, d, J=5.7Hz), 6.73 (1H, d, J=8.2Hz)
6.78-6.85 (4H, m), 7.05 (1H, s), 7.10 (1H, s), 7.18
(2H, d, J=8.6Hz), 7.25-7.45 (6H, m), 7.72 (1H, d,
J=7Hz), 7.91 (1H, d, J=7Hz), 8.05 (1H, d, J=9.3Hz),
15 8.20 (1H, d, J=7Hz), 8.85 (1H, s)

FAB-MASS : m/z = 1390 (M+Na)

Elemental Analysis Calcd. for $\text{C}_{60}\text{H}_{82}\text{N}_9\text{O}_{24}\text{SNa}\cdot 5\text{H}_2\text{O}$:

C 49.41, H 6.36, N 8.64

Found : C 49.77, H 6.71, N 8.71

20

Example 59

IR (KBr) : 3353.6, 1670.1, 1627.6, 1247.7 cm^{-1}

25 NMR (DMSO- d_6 , δ) : 0.86 (3H, t, J=6.5Hz), 0.97 (3H, d,
J=6.8Hz), 1.01 (3H, d, J=5.4Hz), 1.1-1.55 (12H, m),
1.55-1.95 (5H, m), 2.05-4.7 (4H, m), 3.16 (1H, m),
3.5-4.5 (16H, m), 4.6-5.3 (13H, m), 5.55 (1H, d,
J=5.6Hz), 6.7-6.9 (5H, m), 7.05 (1H, s), 7.1 (1H,
s), 7.15 (1H, d, J=8.5Hz), 7.25-7.5 (6H, m), 7.73
(1H, d, J=8.4Hz), 7.92 (1H, d, J=7Hz), 8.08 (1H, d,
J=8.4Hz), 8.18 (1H, d, J=7Hz), 8.84 (1H, s)
30

FAB-MASS : m/z = 1390 (M+Na)

Example 60

35 NMR (DMSO- d_6 , δ) : 0.85 (3H, t, J=6.6Hz), 0.96 (3H, d,
J=6.6Hz), 1.05 (3H, d, J=5.6Hz), 1.1-1.5 (22H, m),

1.5-2.5 (9H, m), 2.5-3.5 (4H, m), 3.5-4.45 (14H, m), 4.45-5.45 (12H, m), 6.72 (1H, d, J=8.2Hz), 6.79 (1H, s), 6.81 (1H, d, J=8.2Hz), 7.04 (1H, s), 7.05-7.5 (8H, m), 7.9-8.3 (3H, m), 8.84 (1H, s)

5 FAB-MASS : m/z = 1325 (M+Na)

Elemental Analysis Calcd. for $C_{58}H_{89}N_8O_{22}SNa \cdot 6H_2O$:

C 49.35, H 7.14, N 7.94

Found : C 49.33, H 7.04, N 7.87

10 Example 61

IR (KBr) : 3400, 1668.1, 1629.6, 1270.9 cm^{-1}

NMR (DMSO- d_6 , δ) : 0.96 (3H, d, J=6.8Hz), 1.06 (3H, d, J=5.7Hz), 1.1-2.0 (33H, m), 2.1-2.5 (4H, m), 3.20 (3H, s), 3.28 (2H, t, J=6.5Hz), 3.1-3.3 (1H, m), 3.6-4.45 (14H, m), 4.6-5.3 (13H, m), 5.49 (1H, d, J=6.1Hz), 6.70 (1H, s), 6.72 (1H, d, J=8.2Hz), 6.80 (1H, d, J=8.2Hz), 7.03 (1H, s), 7.0-7.1 (1H, m), 7.15 (1H, s), 7.2-7.45 (6H, m), 8.0-8.3 (3H, m), 8.83 (1H, s)

20 FAB-MASS : m/z = 1426 (M+Na)

Elemental Analysis Calcd. for $C_{62}H_{94}N_9O_{24}SNa \cdot 5H_2O$:

C 49.82, H 7.01, N 8.43

Found : C 49.86, H 7.31, N 8.40

25 Example 62

IR (KBr) : 3355.5, 1668.1, 1629.6, 1274.7 cm^{-1}

NMR (DMSO- d_6 , δ) : 0.85 (3H, t, J=6.5Hz), 0.96 (3H, d, J=6.7Hz), 1.04 (3H, d, J=5.9Hz), 1.1-2.6 (34H, m), 3.2 (1H, m), 3.6-4.55 (14H, m), 4.7-5.3 (11H, m), 5.47 (1H, d, J=5.9Hz), 6.72 (1H, d, J=8.1Hz), 6.79 (1H, s), 6.81 (1H, d, J=8.1Hz), 7.05 (1H, s), 7.11 (1H, s), 7.2-7.5 (2H, m), 8.0-8.15 (2H, m), 8.20 (1H, d, J=8.0Hz), 8.84 (1H, s)

35 FAB-MASS : m/z = 1235 (M+Na)

Elemental Analysis Calcd. for $C_{51}H_{81}N_8O_{22}SNa \cdot 7H_2O$:

C 45.73, H 7.15, N 8.37

Found : C 45.55, H 7.24, N 8.23

Example 63

5 IR (KBr) : 3353.6, 1664.3, 1627.6 cm^{-1}

NMR (DMSO- d_6 , δ) : 0.86 (3H, t, $J=6.6\text{Hz}$), 0.95 (3H, d, $J=6.7\text{Hz}$), 1.04 (3H, d, $J=5.7\text{Hz}$), 1.2-2.7 (30H, m), 3.16 (1H, m), 3.6-4.5 (13H, m), 4.7-5.3 (11H, m), 5.51 (1H, d, $J=6.0\text{Hz}$), 5.74 (1H, s), 6.72 (1H, d, $J=8.2\text{Hz}$), 6.75 (1H, s), 6.77 (1H, d, $J=8.2\text{Hz}$), 7.05 (1H, s), 7.2-7.5 (3H, m), 8.0-8.3 (3H, m), 8.85 (1H, s)

FAB-MASS : m/z = 1204 (M+Na)

Elemental Analysis Calcd. for $\text{C}_{50}\text{H}_{77}\text{N}_8\text{O}_{21}\text{SNa}\cdot 5\text{H}_2\text{O}$:

15 C 47.24, H 6.90, N 8.81

Found : C 46.98, H 7.12, N 8.72

Example 64

Major product

20 IR (KBr) : 3400, 1675.8, 1631.5, 1511.9, 1234.2 cm^{-1}

NMR (DMSO- d_6 , δ) : 0.96 (3H, d, $J=6.6\text{Hz}$), 1.05 (3H, d, $J=5.8\text{Hz}$), 1.2-1.6 (10H, m), 1.6-2.1 (5H, m), 2.1-2.7 (4H, m), 3.05-3.2 (4H, m), 3.20 (3H, s), 3.29 (2H, t, $J=6.4\text{Hz}$), 3.3-3.5 (5H, m), 3.6-4.5 (15H, m), 4.7-5.3 (11H, m), 5.50 (1H, d, $J=5.8\text{Hz}$), 6.73 (1H, d, $J=8.2\text{Hz}$), 6.8-7.1 (9H, m), 7.2-7.5 (3H, m), 7.81 (2H, d, $J=8.6\text{Hz}$), 8.08 (1H, d, $J=8.2\text{Hz}$), 8.24 (1H, d, $J=7\text{Hz}$), 8.44 (1H, d, $J=7\text{Hz}$), 8.84 (1H, s)

FAB-MASS : m/z = 1403 (M+Na)

30 Elemental Analysis Calcd. for $\text{C}_{61}\text{H}_{85}\text{N}_{10}\text{O}_{23}\text{SNa}\cdot 9\text{H}_2\text{O}$:

C 47.47, H 6.73, N 9.07

Found : C 47.43, H 7.06, N 9.03

Minor product

35 IR (KBr) : 3350, 1668.1, 1631.5, 1511.9, 1234.2 cm^{-1}

NMR (DMSO- d_6 , δ) : 0.96 (3H, d, $J=6.6\text{Hz}$), 1.07 (3H, d, $J=5.8\text{Hz}$), 1.2-1.5 (6H, m), 1.55-2.1 (7H, m), 2.1-2.65 (4H, m), 3.0-3.6 (9H, m), 3.7-4.5 (15H, m), 4.7-5.6 (14H, m), 5.7-6.0 (1H, m), 6.72 (1H, d, $J=8.0\text{Hz}$), 6.75-7.1 (9H, m), 7.25-7.5 (3H, m), 7.81 (2H, d, $J=8.3\text{Hz}$), 8.08 (1H, d, $J=8.2\text{Hz}$), 8.25 (1H, d, $J=7\text{Hz}$), 8.45 (1H, d, $J=7\text{Hz}$), 8.85 (1H, s)

FAB-MASS : $m/z = 1371$ (M+Na)

Elemental Analysis Calcd. for $\text{C}_{60}\text{H}_{81}\text{N}_{10}\text{O}_{22}\text{SNa}\cdot 8\text{H}_2\text{O}$:

C 48.25, H 6.55, N 9.38

Found : C 48.10, H 6.81, N 9.40

Example 65

IR (KBr) : 3450, 1668.1, 1635.3 cm^{-1}

NMR (DMSO- d_6 , δ) : 0.88 (3H, t, $J=6.5\text{Hz}$), 0.96 (3H, d, $J=6.7\text{Hz}$), 1.06 (3H, d, $J=6\text{Hz}$), 1.2-1.5 (6H, m), 1.6-2.1 (5H, m), 2.1-2.7 (4H, m), 3.1-3.4 (9H, m), 3.6-4.5 (15H, m), 4.7-5.3 (11H, m), 5.49 (1H, d, $J=5.8\text{Hz}$), 6.73 (1H, d, $J=8.2\text{Hz}$), 6.8-7.0 (2H, m), 6.83 (2H, d, $J=9.0\text{Hz}$), 6.94 (2H, d, $J=9.0\text{Hz}$), 7.04 (1H, s), 7.12 (1H, t, $J=8.4\text{Hz}$), 7.2-7.5 (3H, m), 7.65-7.8 (2H, m), 8.09 (1H, d, $J=8.4\text{Hz}$), 8.25 (1H, d, $J=7\text{Hz}$), 8.63 (1H, d, $J=7\text{Hz}$), 8.84 (1H, s)

FAB-MASS : $m/z = 1363$ (M+Na)

Elemental Analysis Calcd. for $\text{C}_{58}\text{H}_{78}\text{FN}_{10}\text{O}_{22}\text{SNa}\cdot 5\text{H}_2\text{O}$:

C 48.67, H 6.20, N 9.79

Found : C 48.83, H 6.15, N 9.74

Example 66

IR (KBr) : 3400, 1668.1, 1635.3, 1510.0, 1240.0 cm^{-1}

NMR (DMSO- d_6 , δ) : 0.88 (3H, t, $J=6.6\text{Hz}$), 1.2-1.5 (6H, m), 1.5-2.05 (5H, m), 2.1-2.65 (4H, m), 3.1-3.3 (9H, m), 3.6-4.5 (15H, m), 4.7-5.3 (11H, m), 5.51 (1H, d, $J=5.8\text{Hz}$), 6.73 (1H, d, $J=8.2\text{Hz}$), 6.8-6.9 (4H, m), 6.94 (2H, d, $J=9.2\text{Hz}$), 7.04 (1H, s), 7.24

(1H, d, J=8.5Hz), 7.15-7.5 (3H, m), 7.86 (1H, dd,
J=8.6 and 2.1Hz), 8.02 (1H, d, J=2.1Hz), 8.04 (1H,
d, J=8.4Hz), 8.23 (1H, d, J=7Hz), 8.70 (1H, d,
J=7Hz), 8.84 (1H, s)

5 FAB-MASS : m/z = 1379 (M+Na)

Elemental Analysis Calcd. for $C_{58}H_{78}ClN_{10}O_{22}SNa \cdot 6H_2O$:

C 47.52, H 6.19, N 9.55

Found : C 47.78, H 6.23, N 9.55

10 Example 67

IR (KBr) : 3400, 1670 cm^{-1}

NMR (DMSO- d_6 , δ) : 0.96 (3H, d, J=6.7Hz), 1.05 (3H, d,
J=5.7Hz), 1.4-2.65 (17H, m), 2.65-3.6 (8H, m), 3.6-
4.5 (15H, m), 4.6-5.3 (11H, m), 5.44 (1H, d,
15 J=6.0Hz), 6.73 (1H, d, J=8.2Hz), 6.81 (1H, s), 6.83
(1H, d, J=8.2Hz), 6.98 (2H, d, J=8.9Hz), 7.05 (1H,
s), 7.2-7.5 (3H, m), 7.80 (2H, d, J=8.9Hz), 8.05
(1H, d, J=8.4Hz), 8.26 (1H, d, J=7Hz), 8.39 (1H, d,
J=7Hz), 8.84 (1H, s)

20 FAB-MASS : m/z = 1229 (M+Na)

Elemental Analysis Calcd. for $C_{52}H_{74}N_{10}O_{21}S \cdot 5H_2O$:

C 48.14, H 6.53, N 10.80

Found : C 48.29, H 6.33, N 10.95

25 Example 68

IR (KBr) : 3400, 1652.7, 1635.3, 1511.9, 1241.9 cm^{-1}

NMR (DMSO- d_6 , δ) : 0.88 (3H, t, J=6.6Hz), 0.97 (3H, d,
J=6.7Hz), 1.09 (3H, d, J=5.7Hz), 1.2-1.5 (6H, m),
1.6-2.0 (5H, m), 2.1-2.6 (4H, m), 3.0-3.3 (5H, m),
30 3.6-4.6 (19H, m), 4.7-5.3 (11H, m), 5.53 (1H, d,
J=5.6Hz), 6.73 (1H, d, J=8.2Hz), 6.75-7.0 (2H, m),
6.83 (2H, d, J=9.2Hz), 6.95 (2H, d, J=9.2Hz), 7.05
(1H, s), 7.12 (1H, s), 7.25-7.5 (2H, m), 7.42 (1H,
d, J=9.5Hz), 7.84 (1H, d, J=9.5Hz), 7.9-8.1 (2H,
35 m), 8.71 (1H, d, J=7Hz), 8.84 (1H, s)

FAB-MASS : $m/z = 1347$ (M+Na)

Elemental Analysis Calcd. for $C_{56}H_{77}N_{12}O_{22}SNa \cdot 7H_2O$:

C 46.34, H 6.32, N 11.58

Found : C 46.38, H 6.18, N 11.36

5

Example 69

NMR (DMSO- d_6 , δ) : 0.88 (3H, t, $J=6.6$ Hz), 0.97 (3H, d, $J=6.7$ Hz), 1.08 (3H, d, $J=5.8$ Hz), 1.2-1.5 (6H, m), 1.6-2.05 (5H, m), 2.1-2.6 (4H, m), 3.0-3.3 (5H, m), 3.4-3.55 (4H, m), 3.7-4.6 (15H, m), 4.7-5.3 (11H, m), 5.52 (1H, d, $J=5.8$ Hz), 6.73 (1H, d, $J=8.1$ Hz), 6.8-6.95 (2H, m), 6.83 (2H, d, $J=9.3$ Hz), 6.95 (2H, d, $J=9.3$ Hz) 7.05 (1H, s), 7.14 (1H, s), 7.3-7.6 (3H, m), 7.84 (1H, d, $J=8.6$ Hz), 7.95-8.1 (2H, m), 8.40 (1H, s), 8.42 (1H, d, $J=7$ Hz), 8.84 (1H, s)

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FAB-MASS : $m/z = 1346$ (M+Na)

Elemental Analysis Calcd. for $C_{57}H_{78}N_{11}O_{22}SNa \cdot 5H_2O$:

C 48.40, H 6.27, N 10.89

Found : C 48.32, H 6.44, N 10.86

20

Example 70

IR (KBr) : 3400, 1668.1, 1629.6, 1511.9 cm^{-1}

NMR (DMSO- d_6 , δ) : 0.96 (3H, d, $J=6.7$ Hz), 1.06 (3H, d, $J=5.7$ Hz), 1.15-1.5 (6H, m), 1.6-2.0 (7H, m), 2.1-2.65 (5H, m), 3.1-3.5 (9H, m), 3.6-4.5 (13H, m), 4.7-5.3 (11H, m), 5.46 (1H, d, $J=5.9$ Hz), 6.73 (1H, d, $J=8.2$ Hz), 6.81 (1H, s), 6.84 (1H, d, $J=8.2$ Hz), 6.91 (2H, d, $J=8.7$ Hz), 6.95-7.05 (3H, m), 7.09 (2H, d, $J=8.7$ Hz), 7.25-7.5 (3H, m), 7.81 (2H, d, $J=8.8$ Hz), 8.09 (1H, d, $J=7$ Hz), 8.25 (1H, d, $J=7$ Hz), 8.04 (1H, d, $J=7$ Hz), 8.84 (1H, s)

25

30

FAB-MASS : $m/z = 1327$ (M+Na)

Elemental Analysis Calcd. for $C_{58}H_{77}N_{10}O_{21}SNa \cdot 5H_2O$:

C 49.92, H 6.28, N 10.03

Found : C 49.75, H 6.41, N 10.25

35

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Example 71

IR (KBr) : 3350, 1668.1, 1629.6, 1511.9, 1232.3 cm^{-1}

NMR (DMSO- d_6 , δ) : 0.85 (3H, t, $J=6.5\text{Hz}$), 0.96 (3H, d, $J=6.7\text{Hz}$), 1.06 (3H, d, $J=6.0\text{Hz}$), 1.2-1.4 (6H, m), 1.4-1.6 (2H, m), 1.7-2.1 (3H, m), 2.1-2.7 (6H, m), 3.1-3.5 (9H, m), 3.72 (2H, m), 3.8-4.5 (11H, m), 4.7-5.3 (11H, m), 5.47 (1H, d, $J=5.9\text{Hz}$), 6.73 (1H, d, $J=8.2\text{Hz}$), 6.8-6.9 (2H, m), 6.91 (2H, d, $J=8.6\text{Hz}$), 6.95-7.15 (5H, m), 7.25-7.5 (3H, m), 7.81 (2H, d, $J=8.8\text{Hz}$), 8.09 (1H, d, $J=8.4\text{Hz}$), 8.26 (1H, d, $J=7\text{Hz}$), 8.40 (1H, d, $J=7\text{Hz}$), 8.84 (1H, s)

FAB-MASS : m/z = 1329 (M+Na)

Elemental Analysis Calcd. for $\text{C}_{58}\text{H}_{79}\text{N}_{10}\text{NaO}_{21}\text{S}\cdot 6\text{H}_2\text{O}$:

C 49.22, H 6.48, N 9.90

Found : C 49.33, H 6.67, N 9.89

Example 72

IR (KBr) : 3450, 1668.1, 1631.5, 1240.0 cm^{-1}

NMR (DMSO- d_6 , δ) : 0.96 (3H, d, $J=6.6\text{Hz}$), 1.05 (3H, d, $J=5.6\text{Hz}$), 1.3-1.7 (4H, m), 1.7-2.1 (7H, m), 2.1-2.73 (6H, m), 2.75-3.05 (4H, m), 3.05-4.5 (18H, m), 4.7-5.5 (12H, m), 6.72 (1H, d, $J=8.3\text{Hz}$), 6.77-6.9 (2H, m), 6.96 (2H, d, $J=8.6\text{Hz}$), 7.05 (1H, s), 7.1-7.5 (8H, m), 7.80 (2H, d, $J=8.6\text{Hz}$), 8.06 (1H, d, $J=8.4\text{Hz}$), 8.28 (1H, d, $J=7\text{Hz}$), 8.41 (1H, d, $J=7\text{Hz}$), 8.84 (1H, s)

FAB-MASS : m/z = 1305 (M+Na)

Elemental Analysis Calcd. for $\text{C}_{58}\text{H}_{78}\text{N}_{10}\text{O}_{21}\text{S}\cdot 8\text{H}_2\text{O}$:

C 48.80, H 6.64, N 9.81

Found : C 48.88, H 6.50, N 9.81

Example 73

IR (KBr) : 1673.9, 1646.9, 1510.0 1238.1 cm^{-1}

NMR (DMSO- d_6 , δ) : 0.87 (3H, t, $J=6.4\text{Hz}$), 0.96 (3H, d, $J=6.6\text{Hz}$), 1.05 (3H, d, $J=5.6\text{Hz}$), 1.2-1.5 (6H, m),

1.5-2.0 (9H, m), 2.1-2.8 (11H, m), 3.1-3.4 (5H, m),
3.4-4.5 (17H, m), 4.6-5.5 (12H, m), 6.6-7.0 (9H,
m), 7.04 (1H, s), 7.2-7.5 (3H, m), 7.78 (2H, d,
J=8.7Hz), 8.05 (1H, d, J=8.4Hz), 8.24 (1H, d,
J=7Hz), 8.39 (1H, d, J=7Hz), 8.84 (1H, s)

FAB-MASS : $m/z = 1326$ ($M^+ - SO_3 + Na$)

Elemental Analysis Calcd. for $C_{63}H_{89}N_{11}O_{22}S \cdot 9H_2O$:

C 48.92, H 6.97, N 9.96

Found : C 48.77, H 6.73, N 9.94

Example 74

IR (KBr) : 3450, 1670.1, 1631.5, 1280.5 cm^{-1}

NMR (DMSO- d_6 , δ) : 0.87 (3H, t, J=7.0Hz), 0.96 (3H, t,
J=6.8Hz), 1.05 (3H, d, J=5.6Hz), 1.1-1.65 (13H, m),
1.65-2.1 (7H, m), 2.1-2.65 (5H, m), 3.17 (1H, m),
3.6-4.5 (13H, m), 4.7-5.3 (11H, m), 5.49 (1H, d,
J=5.9Hz), 6.72 (1H, d, J=8.2Hz), 6.82 (1H, d,
J=8.2Hz), 6.84 (1H, s), 7.04 (1H, s), 7.29 (2H, d,
J=8.3Hz), 7.2-7.5 (3H, m), 7.80 (2H, d, J=8.3Hz),
8.10 (1H, d, J=8.4Hz), 8.26 (1H, d, J=7Hz), 8.65
(1H, d, J=7Hz), 8.84 (1H, s)

FAB-MASS : $m/z = 1237$ ($M + Na$)

Elemental Analysis Calcd. for $C_{53}H_{75}N_8O_{21}SNa \cdot 6H_2O$:

C 48.10, H 6.63, N 8.47

Found : C 48.26, H 6.62, N 8.46

Example 75

IR (KBr) : 3400, 1670.1, 1627.6, 1272.8 cm^{-1}

NMR (DMSO- d_6 , δ) : 0.96 (3H, d, J=3.3Hz), 1.08 (3H, d,
J=5.7Hz), 1.2-1.6 (10H, m), 1.6-2.1 (5H, m), 2.1-
2.7 (4H, m), 3.0-3.3 (1H, m), 3.20 (3H, s), 3.29
(2H, t, J=6.4Hz), 3.73 (2H, m), 3.9-4.6 (13H, m),
4.7-5.3 (11H, m), 5.53 (1H, d, J=5.8Hz), 6.73 (1H,
d, J=8.3Hz), 6.83 (1H, d, J=8.3Hz), 6.91 (1H, s),
7.05 (1H, s), 7.23 (1H, dd, J=9.0 and 2.3Hz), 7.3-

7.5 (4H, m), 7.8-8.0 (3H, m), 8.09 (1H, d, J=8.4Hz), 8.33 (1H, d, J=7Hz), 8.44 (1H, s), 8.80 (1H, d, J=7Hz), 8.85 (1H, s)

FAB-MASS : $m/z = 1293$ (M+Na)

5 Elemental Analysis Calcd. for $C_{55}H_{75}N_8O_{23}SNa \cdot 6H_2O$:
C 47.89, H 6.36, N 8.12
Found : C 47.81, H 6.26, N 8.05

Example 76

10 IR (KBr) : 3361.3, 1668.1, 1635.3, 1627.6 cm^{-1}
NMR (DMSO- d_6 , δ) : 0.86 (3H, t, J=6.7Hz), 0.96 (3H, d, J=6.7Hz), 1.09 (3H, d, J=5.8Hz), 1.19-1.25 (8H, m), 1.25-2.00 (5H, m), 2.02-2.53 (4H, m), 2.44 (3H, s), 2.61 (2H, t, J=7.6Hz), 3.05-3.27 (1H, m), 3.55-4.50
15 (13H, m), 4.65-5.65 (12H, m), 6.42 (1H, s), 6.65-6.95 (3H, m), 7.05 (1H, d, J=0.4Hz), 7.13-7.50 (5H, m), 7.50-7.88 (6H, m), 8.10 (1H, d, J=9.0Hz), 8.25 (1H, d, J=8.4Hz), 8.40 (1H, d, J=7.0Hz), 8.85 (1H, s)

20 FAB-MASS : $m/z = 1299.3$ (M+Na-1)
Elemental Analysis Calcd. for $C_{58}H_{77}N_8NaO_{21}S \cdot 5H_2O$:
C 50.94, H 6.41, N 8.19
Found : C 50.99, H 6.40, N 8.15

25 Example 77

IR (Nujol) : 3351.7, 1670.1, 1652.7, 1623.8 cm^{-1}
NMR (DMSO- d_6 , δ) : 0.86 (3H, t, J=6.7Hz), 0.96 (3H, d, J=6.7Hz), 1.06 (3H, d, J=5.8Hz), 1.13-1.45 (8H, m), 1.47-1.96 (5H, m), 2.06-2.66 (8H, m), 2.81 (2H, t, J=7.6Hz), 3.04-3.30 (1H, m), 3.53-4.50 (13H, m), 4.53-5.70 (12H, m), 6.64-6.88 (3H, m), 7.04 (1H, d, J=0.4Hz), 7.13-7.60 (11H, m), 8.10 (1H, d, J=9.0Hz), 8.18 (1H, d, J=8.4Hz), 8.30 (1H, d, J=7.0Hz), 8.85 (1H, s)
30
35 FAB-MASS : $m/z = 1287.4$ (M+Na-1)

Elemental Analysis Calcd. for $C_{57}H_{77}N_8NaO_{21}S \cdot 5H_2O$:

C 50.51, H 6.46, N 8.27

Found : C 50.84, H 6.60, N 8.33

5 Example 78

IR (KBr) : 3361.3, 1683.6, 1670.1, 1662.3, 1652.7,
1646.9, 1635.3, 1627.6, 1623.8 cm^{-1}

10 NMR (DMSO- d_6 , δ) : 0.97 (3H, d, $J=6.7Hz$), 1.07 (3H, d,
 $J=5.6Hz$), 1.28-2.00 (13H, m), 2.08-2.60 (4H, m),
3.07-3.30 (1H, m), 3.60-4.66 (17H, m), 4.66-5.12
(9H, m), 5.11 (1H, d, $J=3.1Hz$), 5.25 (1H, d,
 $J=4.6Hz$), 5.52 (1H, d, $J=6.0Hz$), 6.62-6.95 (4H, m),
6.95-7.15 (3H, m), 7.20-7.50 (3H, m), 7.50-7.85
(7H, m), 8.12 (1H, d, $J=8.4Hz$), 8.35 (1H, d,
15 $J=7.7Hz$), 8.53 (1H, d, $J=7.0Hz$), 8.85 (1H, s)

FAB-MASS : $m/z = 1319.7$ ($M+Na-1$)

Elemental Analysis Calcd. for $C_{57}H_{74}N_8NaO_{22}SF \cdot 8H_2O$:

C 47.49, H 6.29, N 7.77

Found : C 47.79, H 6.16, N 7.93

20 Example 79

IR (KBr) : 3354.9, 1668.1, 1662.3, 1654.6, 1646.9,
1627.6 cm^{-1}

25 NMR (DMSO- d_6 , δ) : 0.85 (3H, t, $J=6.7Hz$), 0.90-1.10
(6H, m), 1.10-1.40 (8H, m), 1.48-1.95 (5H, m),
2.05-2.46 (4H, m), 2.60 (2H, t, $J=7.6Hz$), 3.07-3.23
(1H, m), 3.55-4.45 (14H, m), 4.67-5.32 (11H, m),
5.48-5.63 (1H, m), 6.22 (1H, , $J=5.3Hz$), 6.65-6.89
(3H, m), 6.97-7.15 (2H, m), 7.20-7.68 (10H, m),
30 7.85-8.20 (3H, m), 8.84 (1H, s)

FAB-MASS : $m/z = 1289.4$ ($M+Na-1$)

Elemental Analysis Calcd. for $C_{56}H_{75}N_8NaO_{22}S \cdot 3H_2O$:

C 50.90, H 6.18, N 8.48

Found : C 50.80, H 6.44, N 8.29

Example 80

IR (KBr) : 3361.3, 1664.3, 1631.5, 1600.6 cm^{-1}

NMR (DMSO- d_6 , δ) : 0.86 (3H, t, $J=6.7\text{Hz}$), 0.98 (3H, d, $J=6.7\text{Hz}$), 1.16 (3H, t, $J=5.9\text{Hz}$), 1.20-1.45 (8H, m), 1.50-1.70 (2H, m), 1.70-2.05 (3H, m), 2.10-2.57 (4H, m), 2.63 (2H, t, $J=7.6\text{Hz}$), 3.10-3.30 (1H, m), 3.68-4.50 (13H, m), 4.78-5.32 (11H, m), 5.66 (1H, d, $J=5.7\text{Hz}$), 6.68-7.02 (3H, m), 7.04 (1H, d, $J=0.4\text{Hz}$), 7.25-7.48 (4H, m), 7.60-8.08 (7H, m), 8.10 (1H, d, $J=8.4\text{Hz}$), 8.28 (1H, d, $J=7.7\text{Hz}$), 8.85 (1H, s), 9.30 (1H, d, $J=7.1\text{Hz}$)

FAB-MASS : $m/z = 1287.5$ ($M+\text{Na}-1$)

Elemental Analysis Calcd. for $\text{C}_{55}\text{H}_{73}\text{N}_8\text{NaO}_{22}\text{S}\cdot 3\text{H}_2\text{O}$:

C 50.53, H 6.09, N 8.57

Found : C 50.66, H 6.01, N 8.22

Example 81

IR (KBr) : 3349.7, 1668.1, 1627.6 cm^{-1}

NMR (DMSO- d_6 , δ) : 0.85 (3H, t, $J=6.7\text{Hz}$), 0.96 (3H, d, $J=6.7\text{Hz}$), 1.09 (3H, d, $J=5.8\text{Hz}$), 1.18-1.48 (8H, m), 1.50-2.10 (5H, m), 2.10-2.45 (3H, m), 2.50-2.65 (1H, m), 2.77 (2H, t, $J=7.6\text{Hz}$), 3.05-3.25 (1H, m), 3.60-4.65 (13H, m), 4.67-5.60 (12H, m), 6.65-6.97 (3H, m), 7.05 (1H, d, $J=0.4\text{Hz}$), 7.21-7.43 (4H, m), 7.76 (1H, s), 7.83-8.05 (3H, m), 8.10 (1H, d, $J=9.0\text{Hz}$), 8.29 (1H, d, $J=8.4\text{Hz}$), 8.48 (1H, s), 8.64-9.03 (2H, m)

FAB-MASS : $m/z = 1233.0$ ($M+\text{Na}-1$)

Elemental Analysis Calcd. for $\text{C}_{53}\text{H}_{71}\text{N}_8\text{NaO}_{20}\text{S}\cdot 3\text{H}_2\text{O}$:

C 50.96, H 6.22, N 8.96

Found : C 50.62, H 6.40, N 8.92

Example 82

IR (KBr) : 3361.3, 1670.1, 1627.6 cm^{-1}

NMR (DMSO- d_6 , δ) : 0.88 (3H, t, $J=6.7\text{Hz}$), 0.96 (3H, d,

J=6.7Hz), 1.09 (3H, d, J=5.9Hz), 1.18-1.43 (6H, m),
1.50-2.10 (5H, m), 2.10-2.69 (4H, m), 2.77 (2H, t,
J=7.6Hz), 3.07-3.29 (1H, m), 3.60-4.62 (13H, m),
4.69-5.23 (10H, m), 5.27 (1H, d, J=4.5Hz), 5.55
(1H, d, J=5.9Hz), 6.68-7.00 (3H, m), 7.05 (1H, d,
J=0.4Hz), 7.25-7.53 (4H, m), 7.76 (1H, s), 7.84-
8.05 (3H, m), 8.13 (1H, d, J=8.4Hz), 8.33 (1H, d,
J=7.7Hz), 8.48 (1H, s), 8.73-9.00 (2H, m)

FAB-MASS : $m/z = 1219.4$ (M+Na-1)

Elemental Analysis Calcd. for $C_{52}H_{69}N_8NaO_{21}S \cdot 5H_2O$:

C 48.51, H 6.19, N 8.71

Found : C 48.67, H 6.34, N 8.74

Example 83

IR (KBr) : 3357.5, 1668.1, 1627.6 cm^{-1}

NMR (DMSO- d_6 , δ) : 0.97 (3H, d, J=6.7Hz), 1.07 (3H, d,
J=6.0Hz), 1.20-1.62 (10H, m), 1.62-2.00 (5H, m),
2.10-2.65 (4H, m), 3.20 (3H, s), 3.08-3.45 (1H, m),
3.28 (2H, t, J=6.5Hz), 3.53-4.50 (15H, m), 4.68-
5.13 (9H, m), 5.17 (1H, d, J=3.1Hz), 5.25 (1H, d,
J=4.4Hz), 5.53 (1H, d, J=6.0Hz), 6.68-6.95 (4H, m),
6.95-7.11 (3H, m), 7.20-7.52 (3H, m), 7.55-7.95
(7H, m), 8.13 (1H, d, J=8.4Hz), 8.30 (1H, d,
J=7.7Hz), 8.52 (1H, d, J=7.0Hz), 8.85 (1H, s)

FAB-MASS : $m/z = 1345.2$ (M+Na-1)

Elemental Analysis Calcd. for $C_{59}H_{79}N_8NaO_{23}S \cdot 8H_2O$:

C 48.29, H 6.53, N 7.64

Found : C 48.44, H 6.58, N 7.75

Example 84

IR (KBr) : 3353.6, 1662.3, 1627.6 cm^{-1}

NMR (DMSO- d_6 , δ) : 0.96 (3H, d, J=6.7Hz), 1.07 (3H, d,
J=5.5Hz), 1.40-1.65 (2H, m), 1.65-2.00 (5H, m),
2.00-2.67 (6H, m), 3.08-3.30 (1H, m), 3.50-4.50
(15H, m), 4.68-5.13 (11H, m), 5.18 (1H, d,

J=3.1Hz), 5.26 (1H, d, J=4.5Hz), 5.53 (1H, d,
J=6.0Hz), 5.70-6.00 (1H, m), 6.63-6.95 (4H, m),
6.95-7.13 (3H, m), 7.20-7.52 (3H, m), 7.52-7.95
(7H, m), 8.12 (1H, d, J=8.4Hz), 8.31 (1H, d,
J=7.7Hz), 8.53 (1H, d, J=7.0Hz), 8.85 (1H, s)

FAB-MASS : m/z = 1285.4 (M+Na-1)

Elemental Analysis Calcd. for $C_{56}H_{71}N_8O_{22}SNa \cdot 8H_2O$:

C 47.79, H 6.23, N 7.96

Found : C 47.59, H 6.32, N 8.06

Example 85

IR (KBr) : 3363.2, 1670.1, 1627.6 cm^{-1}

NMR (DMSO- d_6 , δ) : 0.89 (6H, d, J=6.5Hz), 0.96 (3H,
d, J=6.7Hz), 1.07 (3H, d, J=5.7Hz), 1.22-1.41 (2H,
m), 1.50-1.97 (6H, m), 2.11-2.65 (4H, m), 3.10-3.30
(1H, m), 3.60-4.50 (15H, m), 4.70-5.08 (8H, m),
5.10 (1H, d, J=5.6Hz), 5.16 (1H, d, J=3.1Hz), 5.25
(1H, d, J=4.5Hz), 5.50 (1H, d, J=5.9Hz), 6.65-6.92
(4H, m), 6.92-7.12 (3H, m), 7.21-7.50 (3H, m),
7.52-7.90 (7H, m), 8.12 (1H, d, J=8.4Hz), 8.30 (1H,
d, J=7.7Hz), 8.56 (1H, d, J=7.0Hz), 8.85 (1H, s)

FAB-MASS : m/z = 1287.6 (M+Na-1)

Elemental Analysis Calcd. for $C_{56}H_{73}N_8NaO_{22}S \cdot 6.5H_2O$:

C 48.66, H 6.27, N 8.11

Found : C 48.67, H 6.32, N 8.20

Example 86

IR (KBr) : 3361.3, 1683.6, 1670.1, 1654.6, 1635.3,
1623.8 cm^{-1}

NMR (DMSO- d_6 , δ) : 0.97 (3H, d, J=6.7Hz), 1.07 (3H, d,
J=5.6Hz), 1.30-2.00 (11H, m), 2.10-2.70 (4H, m),
3.05-3.15 (1H, m), 3.55-4.70 (17H, m), 4.70-5.11
(9H, m), 5.16 (1H, d, J=3.1Hz), 5.25 (1H, d,
J=4.5Hz), 5.52 (1H, d, J=6.0Hz), 6.65-6.95 (4H, m),
6.95-7.10 (3H, m), 7.10-7.50 (3H, m), 7.50-7.85

(7H, m), 8.12 (1H, d, J=8.4Hz), 8.30 (1H, d,
J=8.3Hz), 8.52 (1H, d, J=7.0Hz), 8.85 (1H, s)

FAB-MASS : $m/z = 1305.2$ (M+Na-1)

Elemental Analysis Calcd. for $C_{56}H_{72}N_8NaO_{22}SF \cdot 6H_2O$:

5 C 48.34, H 6.09, N 8.05

Found : C 48.47, H 6.29, N 7.95

Example 87

IR (KBr) : 3359.4, 1668.1, 1654.6, 1625.7 cm^{-1}

10 NMR (DMSO- d_6 , δ) : 0.97 (3H, d, J=6.7Hz), 1.07 (3H, d,
J=6.0Hz), 1.22-1.62 (6H, m), 1.62-2.00 (5H, m),
2.10-2.65 (4H, m), 3.20 (3H, s), 3.05-3.40 (1H, m),
3.31 (2H, t, J=6.5Hz), 3.60-4.55 (15H, m), 4.65-
5.13 (9H, m), 5.16 (1H, d, J=3.1Hz), 5.26 (1H, d,
15 J=4.4Hz), 5.53 (1H, d, J=6.0Hz), 6.68-6.95 (4H, m),
6.95-7.20 (3H, m), 7.20-7.58 (3H, m), 7.58-7.90
(7H, m), 8.13 (1H, d, J=8.4Hz), 8.32 (1H, d,
J=7.7Hz), 8.53 (1H, d, J=7.0Hz), 8.85 (1H, s)

FAB-MASS : $m/z = 1317.6$ (M+Na-1)

20 Elemental Analysis Calcd. for $C_{57}H_{75}N_8NaO_{23}S \cdot 7H_2O$:

C 48.16, H 6.31, N 7.88

Found : C 48.21, H 6.60, N 7.78

Example 88

25 IR (KBr) : 3350, 2954, 1668, 1629, 1538, 1511, 1454,
1249 cm^{-1}

NMR (DMSO- d_6 , δ) : 0.88 (3H, t, J=7.1Hz), 0.96 (3H, d,
J=7.5Hz), 1.08 (2H, d, J=5.7Hz), 1.2-1.5 (6H, m),
1.6-2.4 (8H, m), 2.6-2.7 (1H, m), 3.1-3.3 (1H, m),
30 3.6-4.5 (19H, m), 4.7-5.3 (8H, m), 6.73 (1H, d,
J=8.2Hz), 6.8-7.1 (5H, m), 7.19 (1H, s), 7.3-7.5
(3H, m), 7.75 (2H, d, J=8.7Hz), 7.8-8.0 (4H, m),
8.08 (1H, d, J=8.9Hz), 8.30 (1H, d, J=7.7Hz), 8.7-
9.0 (3H, m)

35 FAB-MASS : $m/z = 1327$ (M+Na⁺)

Elemental Analysis Calcd. for $C_{57}H_{73}N_{10}O_{22}NaS \cdot 9H_2O$:
C 46.65, H 6.25, N 9.54
Found : C 46.95, H 6.22, N 9.55

5 Example 89

IR (KBr) : 3376, 2931, 2858, 1662, 1631, 1521, 1444,
1245, 1047 cm^{-1}

10 NMR (DMSO- d_6 , δ) : 0.97 (3H, d, $J=6.7Hz$), 1.09 (3H, d,
 $J=5.9Hz$), 1.3-1.6 (6H, m), 1.7-2.1 (5H, m), 2.2-2.4
(3H, m), 2.5-2.6 (1H, m), 3.21 (3H, s), 3.2-3.4
(3H, m), 3.6-4.5 (16H, m), 4.79 (2H, d, $J=6.0Hz$),
4.9-5.2 (5H, m), 5.10 (1H, d, $J=3.6Hz$), 5.18 (1H,
15 d, $J=3.1Hz$), 5.26 (1H, d, $J=4.5Hz$), 5.53 (1H, d,
 $J=6.0Hz$), 6.73 (1H, d, $J=8.2Hz$), 6.8-7.0 (2H, m),
7.0-7.2 (3H, m), 7.3-7.5 (3H, m), 7.6-7.9 (8H, m),
8.01 (2H, d, $J=8.4Hz$), 8.12 (1H, d, $J=8.4Hz$), 8.31
(1H, d, $J=7.7Hz$), 8.79 (1H, d, $J=7.0Hz$), 8.85 (1H,
s)

20 FAB-MASS : $m/z = 1367 (M+Na^+)$

Elemental Analysis Calcd. for $C_{61}H_{77}N_8O_{23}NaS \cdot 6.5H_2O$:
C 50.10, H 6.20, N 7.66
Found : C 50.09, H 6.17, N 7.62

25 Example 90

IR (KBr) : 3363, 2937, 2869, 1646, 1444, 1255 cm^{-1}

30 NMR (DMSO- d_6 , δ) : 0.97 (3H, d, $J=6.7Hz$), 1.08 (3H, d,
 $J=5.7Hz$), 1.2-1.6 (10H, m), 1.7-2.1 (5H, m), 2.1-
2.4 (3H, m), 2.5-2.7 (1H, m), 3.20 (3H, s), 3.2-3.4
(1H, m), 3.6-4.6 (16H, m), 4.7-5.2 (8H, m), 5.16
(1H, d, $J=3.1Hz$), 5.24 (1H, d, $J=4.5Hz$), 5.54 (1H,
d, $J=5.8Hz$), 6.73 (1H, d, $J=8.2Hz$), 6.8-7.0 (2H,
m), 7.1-7.4 (6H, m), 7.97 (2H, d, $J=8.8Hz$), 8.0-8.4
(6H, m), 8.84 (1H, s), 8.92 (1H, d, $J=7.0Hz$)

35 FAB-MASS : $m/z = 1403.6 (M+Na^+)$

Elemental Analysis Calcd. for $C_{59}H_{77}N_{10}O_{23}NaS_2 \cdot 6H_2O$:

C 47.58, H 6.02, N 9.40

Found : C 47.72, H 6.12, N 9.42

Example 91

5 IR (KBr) : 3350, 1668, 1654, 1625, 1537, 1521, 1245,
1047 cm^{-1}

NMR (DMSO- d_6 , δ) : 0.9-1.1 (6H, m), 1.07 (3H, d,
J=5.7Hz), 1.4-2.0 (7H, m), 2.2-2.5 (3H, m), 2.5-2.6
(1H, m), 3.1-3.3 (1H, m), 3.6-4.5 (16H, m), 4.7-5.1
10 (7H, m), 5.09 (1H, d, J=5.6Hz), 5.16 (1H, d,
J=3.1Hz), 5.25 (1H, d, J=4.4Hz), 5.53 (1H, d,
J=6.0Hz), 6.73 (1H, d, J=8.4Hz), 6.8-7.2 (6H, m)
7.2-7.5 (4H, m), 7.5-7.8 (6H, m), 8.11 (1H, d,
J=8.4Hz), 8.32 (1H, d, J=7.7Hz), 8.54 (1H, d,
15 J=7.0Hz), 8.84 (1H, s)

FAB-MASS : $m/z = 1259$ ($M+Na^+$)

Elemental Analysis Calcd. for $C_{54}H_{69}N_8O_{22}NaS \cdot 8H_2O$:

C 46.95, H 6.20, N 8.11

Found : C 47.20, H 6.23, N 8.28

Example 92

IR (KBr) : 3359, 2929, 2852, 1668, 1650, 1631, 1538,
1515 cm^{-1}

NMR (DMSO- d_6 , δ) : 0.96 (3H, d, J=6.7Hz), 1.09 (3H, d,
J=6.1Hz), 1.2-1.6 (5H, m), 1.6-2.5 (10H, m), 2.5-
2.6 (1H, m), 3.18 (1H, m), 3.7-4.5 (15H, m), 4.8-
5.2 (8H, m), 5.17 (1H, d, J=3.1Hz), 5.26 (1H, d,
J=4.5Hz), 5.55 (1H, d, J=5.9Hz), 6.73 (1H, d,
J=8.1Hz), 6.81 (1H, s), 6.85 (1H, s), 7.05 (1H, s),
30 7.2-7.4 (3H, m), 7.45 (2H, d, J=8.2Hz), 7.96 (2H,
d, J=8.2Hz), 8.0-8.2 (4H, s), 8.2-8.3 (1H, m), 8.85
(1H, s), 8.9-9.0 (1H, d, J=7.0Hz)

FAB-MASS : $m/z = 1327.5$ ($M+Na$) $^+$

Elemental Analysis Calcd. for $C_{56}H_{69}N_{10}O_{21}S_2Na \cdot 6H_2O$:

C 47.59, H 5.78, N 9.91

Found : C 47.89, H 5.76, N 9.93

Example 93

IR (KBr) : 3350, 1654, 1629, 1517, 1249, 1047 cm^{-1}

5 NMR (DMSO- d_6 , δ) : 0.9-1.1 (6H, m), 1.11 (3H, d, $J=5.9\text{Hz}$), 1.6-2.0 (5H, s), 2.1-2.4 (3H, s), 2.6-2.7 (1H, m), 3.1-3.3 (1H, m), 3.6-4.5 (16H, m), 4.7-5.2 (7H, m), 5.10 (1H, d, $J=5.6\text{Hz}$), 5.17 (1H, d, $J=3.1\text{Hz}$), 5.25 (1H, d, $J=4.5\text{Hz}$), 5.55 (1H, d, $J=5.7\text{Hz}$), 6.7-6.9 (3H, m), 7.0-7.5 (6H, m), 7.74 (2H, d, $J=8.8\text{Hz}$), 7.91 (2H, d, $J=8.5\text{Hz}$), 8.1-8.4 (8H, m), 8.84 (1H, s), 8.97 (1H, d, $J=7.0\text{Hz}$)

FAB-MASS : $m/z = 1363.5$ ($M+\text{Na}$) $^+$

Elemental Analysis Calcd. for $\text{C}_{59}\text{H}_{69}\text{N}_{10}\text{O}_{23}\text{SNa}\cdot 5\text{H}_2\text{O}$:

15 C 49.51, H 5.56, N 9.79

Found : C 49.39, H 5.63, N 9.77

Example 94

20 IR (KBr) : 3355, 2929, 2856, 1664, 1631, 1519, 1440, 1282 cm^{-1}

NMR (DMSO- d_6 , δ) : 0.84 (3H, t, $J=6.7\text{Hz}$), 0.96 (3H, d, $J=6.7\text{Hz}$), 1.07 (3H, t, $J=5.8\text{Hz}$), 1.2-1.5 (12H, m), 1.7-2.0 (5H, m), 2.2-2.4 (3H, m), 2.5-2.7 (1H, m), 2.94 (2H, t, $J=7.4\text{Hz}$), 3.1-3.3 (1H, m), 3.6-4.6 (14H, m), 4.8-5.2 (7H, m), 5.10 (1H, d, $J=3.6\text{Hz}$), 5.17 (1H, d, $J=3.1\text{Hz}$), 5.26 (1H, d, $J=4.5\text{Hz}$), 5.55 (1H, d, $J=5.9\text{Hz}$), 6.73 (1H, d, $J=8.2\text{Hz}$), 6.8-7.0 (2H, m), 7.0-7.5 (4H, m), 8.0-8.2 (5H, m), 8.27 (1H, d, $J=7.7\text{Hz}$), 8.85 (1H, s), 8.93 (1H, d, $J=7.0\text{Hz}$)

30 FAB-MASS : $m/z = 1279$ ($M+\text{Na}^+$)

Elemental Analysis Calcd. for $\text{C}_{53}\text{H}_{73}\text{N}_{10}\text{O}_{22}\text{SNa}\cdot 5.5\text{H}_2\text{O}$:

C 46.93, H 6.24, N 10.33

Found : C 46.93, H 6.46, N 10.31

Example 95

IR (KBr) : 3363, 1673, 1648, 1538, 1253 cm^{-1}

NMR (DMSO- d_6 , δ) : 0.92 (3H, t, $J=6.8\text{Hz}$), 0.97 (3H, d, $J=6.8\text{Hz}$), 1.10 (3H, d, $J=5.8\text{Hz}$), 1.2-1.5 (6H, m), 1.7-2.1 (5H, m), 2.1-2.4 (3H, m), 2.5-2.6 (1H, m), 3.1-3.3 (1H, m), 3.6-4.5 (16H, m), 4.7-5.1 (9H, m), 5.16 (1H, d, $J=3.1\text{Hz}$), 5.24 (1H, d, $J=4.5\text{Hz}$), 5.54 (1H, d, $J=5.8\text{Hz}$), 6.73 (1H, d, $J=8.2\text{Hz}$), 6.8-7.4 (8H, m), 8.04 (2H, d, $J=8.8\text{Hz}$), 8.13 (2H, d, $J=8.6\text{Hz}$), 8.2-8.4 (4H, m), 8.84 (1H, s), 8.98 (1H, d, $J=7.0\text{Hz}$)

FAB-MASS : $m/z = 1329.6$ ($M+\text{Na}$) $^+$

Elemental Analysis Calcd. for $\text{C}_{56}\text{H}_{71}\text{N}_{10}\text{O}_{23}\text{SNa}\cdot 7\text{H}_2\text{O}$:

C 46.92, H 5.97, N 9.77

Found : C 46.86, H 5.99, N 9.77

Example 96

IR (KBr) : 3355, 2929, 1666, 1648, 1631, 1515, 1442, 1047 cm^{-1}

NMR (DMSO- d_6 , δ) : 0.87 (3H, t, $J=6.7\text{Hz}$), 0.97 (3H, d, $J=6.7\text{Hz}$), 1.10 (3H, d, $J=5.8\text{Hz}$), 1.2-1.5 (10H, m), 1.7-2.1 (5H, m), 2.1-2.4 (3H, m), 2.5-2.6 (1H, m), 3.1-3.3 (1H, m), 3.6-4.6 (16H, m), 4.79 (2H, d, $J=5.9\text{Hz}$), 4.8-5.2 (5H, m), 5.09 (1H, d, $J=5.5\text{Hz}$), 5.16 (1H, d, $J=3.1\text{Hz}$), 5.23 (1H, d, $J=4.5\text{Hz}$), 5.53 (1H, d, $J=5.9\text{Hz}$), 6.73 (1H, d, $J=8.0\text{Hz}$), 6.8-6.9 (2H, m), 7.0-7.5 (6H, m), 7.97 (2H, d, $J=8.8\text{Hz}$), 8.0-8.3 (6H, m), 8.83 (1H, s), 8.88 (1H, d, $J=7.0\text{Hz}$)

FAB-MASS : $m/z = 1373.5$ ($M+\text{Na}$) $^+$

Elemental Analysis Calcd. for $\text{C}_{58}\text{H}_{75}\text{N}_{10}\text{O}_{22}\text{S}_2\text{Na}\cdot 6\text{H}_2\text{O}$:

C 47.73, H 6.01, N 9.60

Found : C 47.57, H 5.92, N 9.53

Example 97

IR (KBr) : 3361, 2925, 2852, 1668, 1650, 1631, 1538,
1452, 1049 cm^{-1}

5 NMR (DMSO- d_6 , δ) : 0.87 (3H, t, $J=6.9\text{Hz}$), 0.96 (3H, d,
 $J=6.7\text{Hz}$), 1.08 (3H, d, $J=5.7\text{Hz}$), 1.2-1.4 (11H, m),
1.4-1.6 (2H, m), 1.7-2.1 (5H, m), 2.1-2.5 (5H, m),
2.5-2.6 (1H, m), 3.1-3.3 (2H, m), 3.7-4.5 (14H, m),
4.7-5.0 (7H, m), 5.09 (1H, d, $J=5.6\text{Hz}$), 5.16 (1H,
d, $J=3.1\text{Hz}$), 5.25 (1H, d, $J=4.5\text{Hz}$), 5.54 (1H, d,
 $J=5.8\text{Hz}$), 6.73 (1H, d, $J=8.2\text{Hz}$), 6.8-7.0 (2H, d),
10 7.04 (1H, s), 7.2-7.5 (3H, m), 8.03 (4H, s), 8.0-
8.3 (2H, m), 8.84 (1H, s), 8.95 (1H, d, $J=7.0\text{Hz}$)

FAB-MASS : $m/z = 1321.9$ ($\text{M}+\text{Na}$)⁺

Elemental Analysis Calcd. for $\text{C}_{55}\text{H}_{75}\text{N}_{10}\text{O}_{21}\text{S}_2\text{Na}\cdot 5\text{H}_2\text{O}$:

15 C 47.54, H 6.17, N 10.08
Found : C 47.38, H 6.12, N 9.98

Example 98

IR (KBr) : 3374, 2937, 2875, 1658, 1629, 1531, 1436,
1255, 1047 cm^{-1}

20 NMR (DMSO- d_6 , δ) : 0.9-1.11 (6H, m), 1.09 (3H, d,
 $J=5.7\text{Hz}$), 1.2-1.5 (4H, m), 1.7-2.1 (5H, m), 2.2-2.5
(3H, m), 2.6-2.7 (1H, m), 3.2-3.3 (1H, m), 3.6-4.5
(16H, m), 4.80 (2H, d, $J=5.8\text{Hz}$), 4.8-5.2 (5H, m),
5.10 (1H, d, $J=5.5\text{Hz}$), 5.17 (1H, d, $J=3.0\text{Hz}$), 5.24
25 (1H, d, $J=4.5\text{Hz}$), 5.53 (1H, d, $J=5.8\text{Hz}$), 6.73 (1H,
d, $J=8.2\text{Hz}$), 6.8-7.0 (2H, m), 7.06 (1H, s), 7.10
(2H, d, $J=8.9\text{Hz}$), 7.2-7.5 (3H, m), 7.68 (1H, s),
7.86 (2H, d, $J=8.8\text{Hz}$), 8.0-8.4 (6H, m), 8.84 (1H,
s), 8.90 (1H, d, $J=7.0\text{Hz}$)

30 FAB-MASS : $m/z = 1314$ ($\text{M}+\text{Na}^+$)

Elemental Analysis Calcd. for $\text{C}_{56}\text{H}_{70}\text{N}_9\text{O}_{23}\text{NaS}\cdot 6\text{H}_2\text{O}$:

C 48.03, H 5.90, N 9.00
Found : C 47.92, H 5.83, N 8.88

35 Example 99

IR (KBr) : 3345, 1646, 1633, 1531, 1257 cm^{-1}

NMR (DMSO-d_6 , δ) : 0.97 (3H, d, $J=6.7\text{Hz}$), 1.11 (3H, d, $J=5.7\text{Hz}$), 1.2-1.6 (10H, m), 1.7-2.5 (8H, m), 2.6-2.7 (1H, m), 3.21 (3H, s), 3.3-3.4 (1H, m), 3.7-4.6 (16H, m), 4.8-5.2 (8H, m), 5.16 (1H, d, $J=3.1\text{Hz}$), 5.24 (1H, d, $J=4.5\text{Hz}$), 5.55 (1H, d, $J=5.7\text{Hz}$), 6.7-6.9 (3H, m), 7.0-7.5 (6H, m), 8.0-8.3 (8H, m), 8.84 (1H, s), 8.96 (1H, d, $J=7.0\text{Hz}$)

FAB-MASS : $m/z = 1387.7$ ($M+\text{Na}^+$)

Elemental Analysis Calcd. for $\text{C}_{59}\text{H}_{77}\text{N}_{10}\text{O}_{24}\text{NaS}\cdot 6\text{H}_2\text{O}$:

C 48.09, H 6.09, N 9.51

Found : C 47.81, H 5.83, N 9.38

Example 100

IR (KBr) : 3357, 1668, 1631, 1429, 1284, 1047 cm^{-1}

NMR (DMSO-d_6 , δ) : 0.97 (3H, d, $J=6.7\text{Hz}$), 1.09 (3H, d, $J=5.8\text{Hz}$), 1.8-2.4 (6H, m), 2.5-2.6 (1H, m), 3.1-3.2 (1H, m), 3.7-4.6 (14H, m), 4.7-5.2 (7H, m), 5.10 (1H, d, $J=5.5\text{Hz}$), 5.17 (1H, d, $J=3.1\text{Hz}$), 5.24 (1H, d, $J=5.5\text{Hz}$), 5.53 (1H, d, $J=5.8\text{Hz}$), 6.75 (1H, d, $J=8.2\text{Hz}$), 6.8-6.9 (2H, m), 7.05 (1H, s), 7.3-7.6 (9H, m), 7.8-7.9 (4H, m), 8.0-8.2 (5H, m), 8.2-8.3 (1H, m), 8.34 (1H, d, $J=9.3\text{Hz}$), 8.7-8.8 (1H, m), 8.85 (1H, s)

FAB-MASS : $m/z = 1332.7$ ($M+\text{Na}^+$)

Elemental Analysis Calcd. for $\text{C}_{58}\text{H}_{65}\text{N}_{10}\text{O}_{22}\text{SNa}\cdot 8\text{H}_2\text{O}$:

C 47.93, H 5.62, N 9.64

Found : C 47.83, H 5.53, N 9.56

Example 101

IR (KBr) : 3353, 2929, 2856, 1666, 1631, 1612, 1496, 1440, 1259 cm^{-1}

NMR (DMSO-d_6 , δ) : 0.87 (3H, t, $J=6.6\text{Hz}$), 0.97 (3H, d, $J=6.5\text{Hz}$), 1.09 (3H, d, $J=5.9\text{Hz}$), 1.2-1.5 (10H, m), 1.7-2.1 (5H, m), 2.2-2.5 (3H, m), 2.6-2.7 (1H, m),

3.1-3.2 (1H, m), 3.6-4.5 (16H, m), 4.7-5.0 (3H, m),
5.0-5.2 (5H, m), 5.10 (1H, d, J=3.1Hz), 5.26 (1H,
d, J=4.2Hz), 5.56 (1H, d, J=5.5Hz), 6.73 (1H, d,
J=8.1Hz), 6.8-7.0 (2H, m), 7.05 (1H, s), 7.1-7.5
(5H, m), 8.0-8.4 (8H, m), 8.85 (1H, s), 8.95 (1H, d,
J=7.0Hz)

FAB-MASS : $m/z = 1357.3$ ($M+Na^+$)

Elemental Analysis Calcd. for $C_{58}H_{75}N_{10}O_{23}NaS \cdot 7H_2O$:

C 47.67, H 6.14, N 9.58

Found : C 47.63, H 6.42, N 9.52

Example 102

IR (KBr) : 3361, 1670, 1648, 1633, 1540, 1519,
1249 cm^{-1}

NMR (DMSO- d_6 , δ) : 0.89 (3H, t, J=7.0Hz), 0.97 (3H, d,
J=6.8Hz), 1.10 (3H, d, J=5.7Hz), 1.2-1.5 (6H, m),
1.6-2.4 (8H, m), 2.5-2.7 (1H, m), 3.1-3.3 (1H, m),
3.6-4.5 (16H, m), 4.80 (2H, d, J=5.8Hz), 4.8-5.2
(5H, m), 5.10 (1H, d, J=5.4Hz), 5.18 (1H, d,
J=3.1Hz), 5.25 (1H, d, J=4.3Hz), 5.55 (1H, d,
J=5.7Hz), 6.73 (1H, d, J=8.2Hz), 6.8-7.0 (2H, m),
7.0-7.5 (6H, m), 8.02 (1H, d, J=5.3Hz), 8.0-8.4
(4H, m), 8.42 (2H, d, J=8.4Hz), 8.48 (2H, d,
J=8.9Hz), 8.8-9.0 (3H, m)

FAB-MASS : $m/z = 1339.3$ ($M+Na^+$)

Elemental Analysis Calcd. for $C_{58}H_{73}N_{10}O_{22}SNa \cdot 6H_2O$:

C 48.87, H 6.01, N 9.83

Found : C 49.16, H 5.92, N 9.86

Example 103

IR (KBr) : 3350, 2971, 2859, 1672, 1629, 1537, 1442,
1247, 1047 cm^{-1}

NMR (DMSO- d_6 , δ) : 0.96 (3H, d, J=6.8Hz), 1.0-1.2 (6H,
m), 1.2-1.6 (12H, m), 1.7-2.5 (8H, m), 2.5-2.6 (1H,
m), 3.2-3.6 (7H, m), 3.7-4.5 (16H, m), 4.76 (2H, d,

J=5.6Hz), 4.8-5.1 (5H, m), 5.09 (1H, d, J=5.5Hz),
5.16 (1H, d, J=3.1Hz), 5.23 (1H, d, J=5.5Hz), 5.51
(1H, d, J=5.9Hz), 6.73 (1H, d, J=8.2Hz), 6.8-6.9
(2H, m), 7.0-7.1 (3H, m), 7.3-7.5 (3H, m), 7.67
(2H, d, J=6.9Hz), 7.71 (2H, d, J=6.9Hz), 7.95 (2H,
d, J=8.4Hz), 8.05 (1H, d, J=7.0Hz), 8.23 (1H, d,
J=7.7Hz), 8.70 (1H, d, J=7.0Hz), 8.84 (1H, s)

FAB-MASS : $m/z = 1377.1$ ($M+Na^+$)

Elemental Analysis Calcd. for $C_{60}H_{83}N_8O_{24}NaS \cdot 5H_2O$:

C 49.86, H 6.49, N 7.75

Found : C 49.74, H 6.73, N 7.68

Example 104

IR (KBr) : 3349, 2937, 2858, 1672, 1629, 1537, 1444,
1249, 1047 cm^{-1}

NMR (DMSO- d_6 , δ) : 0.96 (3H, d, J=6.7Hz), 1.08 (3H, d,
J=5.6Hz), 1.2-1.7 (14H, m), 1.7-2.1 (5H, m), 2.1-
2.4 (5H, m), 2.5-2.6 (1H, m), 3.1-3.2 (1H, m), 3.4-
3.6 (4H, m), 3.7-4.5 (16H, m), 4.77 (2H, d,
J=5.7Hz), 4.8-5.2 (5H, m), 5.09 (1H, d, J=5.6Hz),
5.16 (1H, d, J=3.1Hz), 5.24 (1H, d, J=4.5Hz), 5.51
(1H, d, J=5.8Hz), 6.73 (1H, d, J=8.2Hz), 6.8-6.9
(2H, m), 7.0-7.1 (3H, m), 7.3-7.5 (3H, m), 7.6-7.8
(4H, m), 7.96 (2H, d, J=8.4Hz), 8.10 (1H, d,
J=8.4Hz), 8.24 (1H, d, J=7.7Hz), 8.71 (1H, d,
J=7.0Hz), 8.89 (1H, s)

FAB-MASS : $m/z = 1386.5$ ($M+Na^+$)

Elemental Analysis Calcd. for $C_{61}H_{82}N_9O_{23}NaS \cdot 6H_2O$:

C 49.76, H 6.43, N 8.56

Found : C 49.99, H 6.39, N 8.52

Example 105

IR (KBr) : 3350, 2933, 2856, 1664, 1631, 1604, 1511,
1450, 1243, 1045 cm^{-1}

NMR (DMSO- d_6 , δ) : 0.86 (3H, t, J=6.7Hz), 0.96 (3H, d,

J=6.5Hz), 1.05 (3H, d, J=5.7Hz), 1.2-1.5 (8H, m),
1.6-2.0 (5H, m), 2.1-2.4 (3H, m), 2.5-2.6 (1H, m),
3.0-3.3 (5H, m), 3.6-4.4 (20H, m), 4.7-5.1 (7H, m),
5.10 (1H, d, J=5.5Hz), 5.16 (1H, d, J=3.1Hz), 5.27
(1H, d, J=4.5Hz), 5.51 (1H, d, J=6.0Hz), 6.7-7.1
(9H, m), 7.2-7.5 (3H, m), 8.0-8.2 (2H, m), 8.2-8.4
(1H, m), 8.4-8.6 (1H, m), 8.66 (1H, d, J=2.2Hz),
8.85 (1H, s)

FAB-MASS : $m/z = 1360$ ($M+Na^+$)

Elemental Analysis Calcd. for $C_{58}H_{80}N_{11}O_{22}SNa \cdot 6H_2O$:

C 48.16, H 6.41, N 10.65

Found : C 47.91, H 6.31, N 10.56

Example 106

IR (KBr) : 3369, 3345, 2935, 1672, 1629, 1511, 1245,
1047 cm^{-1}

NMR (DMSO- d_6 , δ) : 0.96 (3H, d, J=6.7Hz), 1.06 (3H, d,
J=5.8Hz), 1.3-1.6 (10H, m), 1.6-2.0 (5H, m), 2.1-
2.4 (3H, m), 2.5-2.6 (1H, m), 3.20 (3H, s), 3.28
(2H, t, J=6.4Hz), 3.1-3.4 (5H, m), 3.7-4.5 (20H,
m), 4.7-5.1 (7H, m), 5.08 (1H, d, J=5.5Hz), 5.15
(1H, d, J=3.1Hz), 5.23 (1H, d, J=4.5Hz), 5.48 (1H,
d, J=5.8Hz), 6.73 (1H, d, J=8.2Hz), 6.82 (2H, d,
J=9.1Hz), 6.94 (2H, d, J=9.1Hz), 6.9-7.0 (1H, m),
7.04 (1H, s), 7.3-7.5 (3H, m), 8.0-8.1 (2H, m),
8.27 (1H, d, J=7.7Hz), 8.49 (1H, d, J=7.0Hz), 8.66
(1H, d, J=2.2Hz), 8.84 (1H, s)

FAB-MASS : $m/z = 1404$ ($M+Na^+$)

Example 107

IR (KBr) : 3357, 1647, 1631, 1537, 1444, 1249,
1049 cm^{-1}

NMR (DMSO- d_6 , δ) : 0.9-1.1 (6H, m), 1.09 (3H, d,
J=5.9Hz), 1.6-2.4 (8H, m), 2.4-2.5 (1H, m), 3.1-3.3
(1H, m), 3.6-4.5 (16H, m), 4.8-5.2 (7H, m), 5.10

(1H, d, J=5.6Hz), 5.17 (1H, d, J=3.1Hz), 5.25 (1H, d, J=4.5Hz), 5.55 (1H, d, J=5.9Hz), 6.73 (1H, d, J=8.2Hz), 6.8-7.0 (2H, m), 7.0-7.6 (6H, m), 7.73 (2H, d, J=8.7Hz), 7.86 (2H, d, J=8.5Hz), 8.0-8.3 (8H, m), 8.84 (1H, s), 8.9-9.0 (1H, m)

FAB-MASS : $m/z = 1379.4$ (M+Na)⁺

Elemental Analysis Calcd. for C₅₉H₆₉N₁₀O₂₂S₂Na·6H₂O :

C 48.36, H 5.57, N 9.56

Found : C 48.18, H 5.60, N 9.36

The Object Compounds (108) to (117) were obtained according to a similar manner to that of Example 27.

Example 108

IR (KBr) : 3350, 2933, 1670, 1627, 1521, 1436, 1272, 1047 cm⁻¹

NMR (DMSO-d₆, δ) : 0.85 (3H, t, J=6.7Hz), 0.92 (3H, d, J=6.7Hz), 1.1-1.4 (11H, m), 1.7-2.4 (9H, m), 3.1-3.2 (1H, m), 3.5-5.4 (27H, m), 6.6-7.2 (8H, m), 7.5-7.8 (3H, m), 7.8-8.0 (3H, m), 8.1-8.8 (3H, m)

FAB-MASS : $m/z = 1249.4$ (M+Na)⁺

Elemental Analysis Calcd. for C₅₂H₇₁N₁₀O₂₁NaS·7H₂O :

C 46.15, H 6.33, N 10.35

Found : C 46.12, H 6.35, N 10.24

Example 109

IR (Kbr pelet) : 3361, 2933, 2856, 1670, 1652, 1616, 1540, 1508, 1448, 1261, 1047 cm⁻¹

NMR (DMSO-d₆, δ) : 0.86 (3H, t, J=6.6Hz), 0.97 (3H, d, J=6.8Hz), 1.12 (3H, d, J=6.8Hz), 1.2-1.5 (10H, m), 1.7-2.0 (5H, m), 2.2-2.6 (4H, m), 3.1-3.2 (1H, m), 3.7-4.4 (16H, m), 4.8-5.3 (10H, m), 5.59 (1H, d, J=6.0Hz), 6.7-6.9 (3H, m), 7.0-7.4 (7H, m), 7.8-8.2 (4H, m), 8.8-9.0 (2H, m)

FAB-MASS : $m/z = 1280.3$ (M+Na)⁺

Elemental Analysis Calcd. for $C_{54}H_{72}N_9O_{23}NaS \cdot 7H_2O$:

C 46.45, H 6.21, N 9.03

Found : C 46.68, H 6.44, N 9.03

5 Example 110

IR (KBr) : 3350, 2931, 1670, 1627, 1540, 1436, 1276,
1047 cm^{-1}

10 NMR (DMSO- d_6 , δ) : 0.87 (3H, t, $J=6.8Hz$), 0.93 (2H, d,
 $J=8.8Hz$), 1.08 (2H, d, $J=5.9Hz$), 1.2-1.4 (4H, m),
1.5-1.7 (2H, m), 1.7-2.1 (3H, m), 2.1-2.4 (3H, m),
2.6-2.7 (3H, m), 3.1-3.3 (1H, m), 3.6-4.5 (17H, m),
4.7-5.4 (8H, m), 6.73 (1H, d, $J=8.2Hz$), 6.83 (2H,
d, $J=8.2Hz$), 7.0-7.1 (1H, m), 7.2-7.5 (5H, m), 7.65
15 (2H, d, $J=8.2Hz$), 7.74 (2H, d, $J=8.4Hz$), 7.98 (2H,
d, $J=8.4Hz$), 8.08 (1H, d, $J=8.5Hz$), 8.25 (1H, d,
 $J=8.5Hz$), 8.74 (1H, d, $J=7.6Hz$), 8.7-9.0 (1H, br)

FAB-MASS : $m/z = 1231.2$ ($M+Na^+$)

Elemental Analysis Calcd. for $C_{53}H_{69}N_8O_{21}NaS \cdot 3H_2O$:

C 50.39, H 5.98, N 8.87

20 Found : C 50.34, H 6.25, N 8.90

Example 111

IR (KBr) : 3353.6, 1670.1, 1652.7, 1623.8 cm^{-1}

25 NMR (DMSO- d_6 , δ) : 0.96 (3H, d, $J=6.7Hz$), 1.07 (3H, d,
 $J=5.6Hz$), 1.20-1.62 (8H, m), 1.62-2.00 (5H, m),
2.10-2.65 (4H, m), 3.20 (3H, s), 3.08-3.40 (1H, m),
3.30 (2H, t, $J=6.5Hz$), 3.53-4.50 (15H, m), 4.68-
5.13 (9H, m), 5.16 (1H, d, $J=2.9Hz$), 5.26 (1H, d,
 $J=4.5Hz$), 5.53 (1H, d, $J=5.9Hz$), 6.68-6.95 (4H, m),
30 6.95-7.11 (3H, m), 7.20-7.52 (3H, m), 7.55-7.95
(7H, m), 8.13 (1H, d, $J=8.4Hz$), 8.31 (1H, d,
 $J=7.7Hz$), 8.53 (1H, d, $J=7.0Hz$), 8.85 (1H, s)

FAB-MASS : $m/z = 1331.5$ ($M+Na-1$)

Elemental Analysis Calcd. for $C_{58}H_{77}N_8NaO_{23}S \cdot 6H_2O$:

35 C 49.15, H 6.33, N 7.91

Found : C 49.07, H 6.53, N 7.84

Example 112

5 IR (KBr) : 3350, 2937, 1673, 1646, 1631, 1538, 1519,
1456, 1247, 1049 cm^{-1}

NMR (DMSO- d_6 , δ) : 0.97 (3H, d, $J=6.6\text{Hz}$), 1.07 (3H, d,
 $J=5.7\text{Hz}$), 1.3-2.4 (25H, m), 2.5-2.6 (1H, m), 3.2-
3.4 (1H, m), 3.5-4.6 (20H, m), 4.8-5.7 (11H, m),
6.73 (1H, d, $J=8.0\text{Hz}$), 6.9-7.0 (2H, m), 7.0-7.2
10 (3H, m), 7.3-7.6 (3H, m), 7.74 (2H, d, $J=8.5\text{Hz}$),
7.77 (2H, d, $J=8.3\text{Hz}$), 8.02 (2H, d, $J=8.3\text{Hz}$), 8.13
(1H, d, $J=8.4\text{Hz}$), 8.30 (1H, d, $J=7.7\text{Hz}$), 8.77 (1H,
d, $J=7.0\text{Hz}$), 8.85 (1H, s)

FAB-MASS : $m/z = 1389$ ($M+\text{Na}^+$)

15 Elemental Analysis Calcd. for $\text{C}_{61}\text{H}_{83}\text{N}_8\text{O}_{24}\text{NaS}\cdot 7\text{H}_2\text{O}$:

C 49.06, H 6.55, N 7.50

Found : C 49.03, H 6.54, N 7.56

Example 113

20 NMR (DMSO- d_6 , δ) : 0.84 (3H, t, $J=6.7\text{Hz}$), 0.96 (3H, d,
 $J=6.7\text{Hz}$), 1.07 (3H, d, $J=5.9\text{Hz}$), 1.1-1.3 (14H, m),
1.7-2.1 (5H, m), 2.2-2.5 (3H, m), 2.6-2.7 (1H, m),
3.1-3.3 (1H, m), 3.7-4.5 (16H, m), 4.7-5.1 (7H, m),
5.10 (1H, d, $J=5.5\text{Hz}$), 5.16 (1H, d, $J=3.1\text{Hz}$), 5.25
25 (1H, d, $J=4.5\text{Hz}$), 5.49 (1H, d, $J=5.7\text{Hz}$), 6.53 (1H,
d, $J=3.1\text{Hz}$), 6.73 (1H, d, $J=8.2\text{Hz}$), 6.8-6.9 (2H,
m), 7.05 (1H, m), 7.31 (1H, d, $J=8.1\text{Hz}$), 7.4-7.6
(4H, m), 7.70 (1H, d, $J=6.7\text{Hz}$), 8.08 (1H, d,
 $J=8.4\text{Hz}$), 8.18 (1H, s), 8.31 (1H, d, $J=7.7\text{Hz}$), 8.57
30 (1H, d, $J=7.0\text{Hz}$), 8.85 (1H, s)

FAB-MASS : $m/z = 1264$ ($M+\text{Na}^+$)

Elemental Analysis Calcd. for $\text{C}_{54}\text{H}_{76}\text{N}_9\text{O}_{21}\text{NaS}\cdot 6\text{H}_2\text{O}$:

C 48.03, H 6.57, N 9.34

Found : C 48.02, H 6.61, N 9.28

Example 114

IR (KBr) : 3350, 2937, 1668, 1631, 1537, 1247, 1047 cm^{-1}

NMR (DMSO- d_6 , δ) : 0.85 (3H, t, $J=7.4\text{Hz}$), 0.96 (3H, d, $J=6.5\text{Hz}$), 1.07 (3H, d, $J=5.7\text{Hz}$), 1.3-1.7 (7H, m), 1.7-2.1 (5H, m), 2.2-2.4 (3H, m), 2.6-2.7 (1H, m), 3.0-3.8 (16H, m), 3.8-4.6 (11H, m), 4.7-5.3 (6H, m), 6.73 (1H, d, $J=8.2\text{Hz}$), 6.8-7.0 (2H, m), 7.0-7.2 (3H, m), 7.3-7.5 (3H, m), 7.6-7.8 (4H, m), 7.96 (2H, d, $J=8.3\text{Hz}$), 8.11 (1H, d, $J=8.2\text{Hz}$), 8.26 (1H, d, $J=7.6\text{Hz}$), 8.6-9.0 (2H, m)

FAB-MASS : $m/z = 1319.4$ ($M+\text{Na}^+$)

Elemental Analysis Calcd. for $\text{C}_{57}\text{H}_{77}\text{N}_8\text{O}_{23}\text{NaS}\cdot 8\text{H}_2\text{O}$:

C 47.50, H 6.50, N 7.77

Found : C 47.72, H 6.85, N 7.85

Example 115

IR (KBr) : 3350, 1666, 1631, 1546, 1276, 1247 cm^{-1}

NMR (DMSO- d_6 , δ) : 0.97 (3H, d, $J=7.5\text{Hz}$), 1.08 (3H, d, $J=5.7\text{Hz}$), 1.4-1.6 (4H, m), 1.6-2.1 (5H, m), 2.1-2.4 (3H, m), 2.5-2.6 (1H, m), 3.1-3.3 (1H, m), 3.23 (3H, s), 3.3-3.5 (2H, m), 3.7-4.5 (16H, m), 4.79 (2H, d, $J=6.2\text{Hz}$), 4.8-5.1 (5H, m), 5.11 (1H, d, $J=5.6\text{Hz}$), 5.18 (1H, d, $J=3.1\text{Hz}$), 5.26 (1H, d, $J=4.4\text{Hz}$), 5.54 (1H, d, $J=5.7\text{Hz}$), 6.73 (1H, d, $J=8.1\text{Hz}$), 6.8-7.0 (2H, m), 7.0-7.1 (3H, m), 7.3-7.5 (3H, m), 7.6-7.9 (8H, m), 8.01 (2H, d, $J=8.4\text{Hz}$), 8.08 (1H, d, $J=8.4\text{Hz}$), 8.32 (1H, d, $J=7.7\text{Hz}$), 8.80 (1H, d, $J=7.0\text{Hz}$), 8.85 (1H, s)

FAB-MASS : $m/z = 1353.9$ ($M+\text{Na}^+$)

Elemental Analysis Calcd. for $\text{C}_{60}\text{H}_{75}\text{N}_8\text{O}_{23}\text{NaS}\cdot 9.5\text{H}_2\text{O}$:

C 47.96, H 6.31, N 7.46

Found : C 47.97, H 6.25, N 7.41

Example 116

IR (KBr) : 3450, 2935, 1675, 1650, 1540, 1513, 1454,

1047 cm^{-1}

NMR (DMSO-d_6 , δ) : 0.97 (3H, d, $J=6.7\text{Hz}$), 1.09 (3H, d, $J=5.9\text{Hz}$), 1.60 (6H, s), 1.7-2.4 (6H, m), 2.5-2.6 (1H, m), 3.1-3.6 (5H, m), 3.7-4.5 (14H, m), 4.7-5.0 (3H, m), 5.0-5.2 (4H, m), 5.11 (1H, d, $J=5.5\text{Hz}$), 5.18 (1H, d, $J=3.1\text{Hz}$), 5.26 (1H, d, $J=4.5\text{Hz}$), 5.56 (1H, d, $J=6.0\text{Hz}$), 6.8-7.5 (9H, m), 7.84 (2H, d, $J=8.8\text{Hz}$), 8.0-8.4 (6H, m), 8.85 (1H, s), 8.91 (1H, d, $J=7.0\text{Hz}$)

FAB-MASS : $m/z = 1328$ ($\text{M}+\text{Na}$)⁺

Elemental Analysis Calcd. for $\text{C}_{55}\text{H}_{68}\text{N}_{11}\text{O}_{21}\text{S}_2\text{Na}\cdot 8\text{H}_2\text{O}$:
C 45.55, H 5.84, N 10.62
Found : C 45.62, H 5.70, N 10.54

Example 117

IR (KBr) : 3350, 2939, 1664, 1627, 1531, 1446, 1249, 1049 cm^{-1}

NMR (DMSO-d_6 , δ) : 0.8-1.0 (6H, m), 1.4-1.9 (9H, m), 2.0-2.5 (4H, m), 3.1-3.2 (1H, m), 3.22 (3H, s), 3.3-3.4 (2H, m), 3.51 (2H, s), 3.6-4.4 (16H, m), 4.7-5.2 (7H, m), 5.07 (1H, d, $J=5.6\text{Hz}$), 5.17 (1H, d, $J=3.1\text{Hz}$), 5.23 (1H, d, $J=4.5\text{Hz}$), 5.54 (1H, d, $J=5.9\text{Hz}$), 6.7-6.8 (3H, m), 7.0-7.4 (8H, m), 7.5-7.7 (4H, m), 7.70 (4H, s), 8.1-8.2 (2H, m), 8.51 (1H, d, $J=7.0\text{Hz}$), 8.83 (1H, s)

FAB-MASS : $m/z = 1367.6$ ($\text{M}+\text{Na}$)⁺

Elemental Analysis Calcd. for $\text{C}_{61}\text{H}_{77}\text{N}_8\text{O}_{23}\text{SNa}\cdot 6.5\text{H}_2\text{O}$:
C 50.01, H 6.20, N 7.66
Found : C 50.30, H 6.50, N 7.75

Example 118

To a solution of The Object Compound (61) (0.25 g) in methanol (50 ml) was added dry 10% palladium on carbon (0.2 g) and stirred for 6 hours under hydrogen atmosphere. The palladium on carbon was filtered off, and the filtrate was

evaporated under reduced pressure to give Object Compound 118 (179 mg).

IR (KBr) : 3400, 1668.1, 1627.6 cm^{-1}

NMR (DMSO- d_6 , δ) : 0.92 (3H, d, $J=6.7\text{Hz}$), 1.1-2.45

5 (40H, m), 3.20 (3H, s), 3.28 (2H, t, $J=6.5\text{Hz}$), 3.0-3.4 (1H, m), 3.5-4.7 (14H, m), 4.95-5.5 (12H, m), 6.55 (1H, d, $J=8.4\text{Hz}$), 6.84 (1H, s), 6.86 (1H, d, $J=8.4\text{Hz}$), 7.0-7.3 (4H, m), 7.9-8.3 (4H, m)

FAB-MASS : $m/z = 1292$ (M+Na)

10 Elemental Analysis Calcd. for $\text{C}_{54}\text{H}_{88}\text{N}_9\text{O}_{22}\text{SNa}\cdot 5\text{H}_2\text{O}$:

C 47.67, H 7.26, N 9.26

Found : C 47.72, H 7.35, N 8.95

15 The Object Compounds (119) to (121) were obtained according to a similar manner to that of Example 118.

Example 119

NMR (DMSO- d_6 , δ) : 0.87 (3H, t, $J=6.6\text{Hz}$), 1.00 (3H, d, $J=7.3\text{Hz}$), 1.03 (3H, d, $J=6.0\text{Hz}$), 1.2-1.5 (4H, m),
20 1.5-2.0 (5H, m), 2.1-2.7 (8H, m), 3.17 (1H, m), 3.6-4.5 (14H, m), 4.65-5.7 (12H, m), 6.72 (1H, d, $J=8.1\text{Hz}$), 6.75 (1H, s), 6.80 (1H, d, $J=8.1\text{Hz}$), 7.05 (1H, s), 7.1-7.7 (15H, m), 8.0-8.6 (4H, m), 8.85 (1H, s)

25 FAB-MASS : $m/z = 1274$ (M+Na)

Elemental Analysis Calcd. for $\text{C}_{55}\text{H}_{74}\text{N}_9\text{O}_{21}\text{SNa}\cdot 7\text{H}_2\text{O}$:

C 47.93, N 6.43, N 9.15

Found : C 48.12, N 6.56, N 9.03

30 Example 120

IR (KBr) : 3355.5, 1672.0 1629.6 cm^{-1}

NMR (DMSO- d_6 , δ) : 0.86 (3H, t, $J=6.6\text{Hz}$), 0.98 (3H, d, $J=6.5\text{Hz}$), 1.03 (3H, d, $J=6.0\text{Hz}$), 1.2-2.6 (21H, m),
35 3.18 (1H, m), 3.6-4.5 (16H, m), 4.65-5.55 (12H, m), 6.6-7.5 (10H, m), 8.0-8.6 (4H, m), 8.89 (1H, s)

FAB-MASS : $m/z = 1256$ (M+Na)

Example 121

IR (KBr) : 3357.5, 1660.4, 1629.6, 1249.6 cm^{-1}

5 NMR (DMSO- d_6 , δ) : 0.86 (3H, t, $J=6.6\text{Hz}$), 0.96 (3H, d, $J=6.8\text{Hz}$), 1.03 (3H, d, $J=6.0\text{Hz}$), 1.1-1.5 (12H, m),
1.6-2.0 (5H, m), 2.0-2.5 (4H, m), 3.07 (1H, m),
3.5-4.5 (16H, m), 4.6-5.6 (12H, m), 6.72 (1H, d, $J=8.1\text{Hz}$), 6.7-6.9 (4H, m), 7.04 (1H, s), 7.16 (1H,
10 s), 7.1-7.5 (2H, m), 7.25 (2H, d, $J=8.6\text{Hz}$), 8.0-8.2
(3H, m), 8.46 (1H, d, $J=7\text{Hz}$), 8.84 (1H, s)

FAB-MASS : $m/z = 1256$ (M+Na)

Elemental Analysis Calcd. for $\text{C}_{52}\text{H}_{76}\text{N}_9\text{O}_{22}\text{SNa}\cdot 7\text{H}_2\text{O}$:

C 45.91, H 6.67, N 9.27

15 Found : C 45.98, H 6.67, N 9.10

Example 122

A solution of Object Compound (11) (795 mg) in water (16 ml) was left for 240 hours. The solution was subjected to
20 column chromatography on ODS (YMC-gel ODS-AMS50) and eluted
with 25% $\text{CH}_3\text{CN}/\text{H}_2\text{O}$. The fractions containing Object Compound
were combined and the acetonitrile was removed under reduced
pressure. The residue was lyophilized to give Object
Compound (123) (38 mg).

25 IR (KBr) : 3361, 2956, 2875, 1668, 1627, 1521, 1249,
1047 cm^{-1}

NMR (DMSO- d_6 , δ) : 0.8-1.5 (19H, m), 1.6-2.4 (13H, m),
3.1-3.2 (1H, m), 3.5-4.1 (12H, m), 4.1-4.7 (10H,
m), 4.9-5.6 (5H, m), 5.98 (1H, d, $J=10.6\text{Hz}$), 6.36
30 (1H, d, $J=10.6\text{Hz}$), 6.7-7.3 (12H, m), 7.4-8.0 (7H,
m)

FAB-MASS : $m/z = 1273.1$ (M+ Na^+)

Elemental Analysis Calcd. for $\text{C}_{55}\text{H}_{71}\text{N}_8\text{O}_{22}\text{NaS}\cdot 11\text{H}_2\text{O}$:

C 45.58, H 6.47, N 7.73

35 Found : C 45.83, H 6.26, N 7.75

The Object Compound (123) was obtained according to a similar manner to that of Example 118.

Example 123

5 IR (KBr) : 3349.7, 1670.1, 1627.6 cm^{-1}
NMR (DMSO- d_6 , δ) : 0.87 (3H, t, $J=7.2\text{Hz}$), 0.96 (3H, d, $J=6.7\text{Hz}$), 1.13 (3H, d, $J=5.7\text{Hz}$), 1.18-1.55 (10H, m), 1.58-2.08 (5H, m), 2.08-2.90 (4H, m), 2.90-3.30 (2H, m), 3.60-4.50 (17H, m), 4.70-5.70 (12H, m),
10 6.65-7.60 (11H, m), 7.80 (2H, br s), 7.95-8.23 (2H, m), 8.75 (1H, d, $J=7.0\text{Hz}$), 8.85 (1H, s)
FAB-MASS : $m/z = 1114.4$ (M-SO₄-2)
Elemental Analysis Calcd. for C₅₂H₇₇N₉O₂₁S·6H₂O :
C 47.88, H 6.88, N 9.66
15 Found : C 47.60, H 6.74, N 9.53

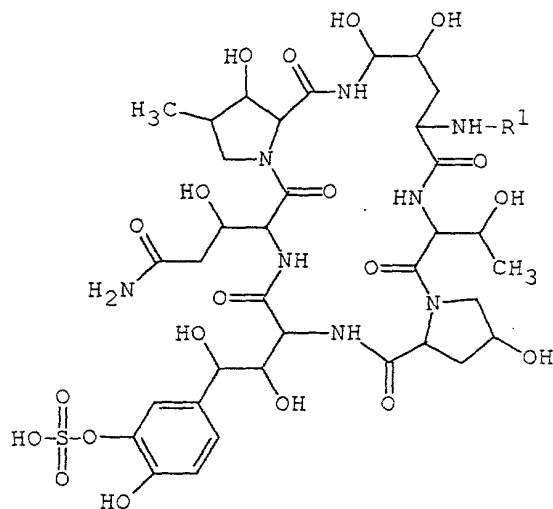
The following compound (124) was obtained according to a similar manner to that of Example 1.

Example 124

20 IR (KBr) : 3324, 2937, 2873, 1664, 1629, 1442, 1257 cm^{-1}
NMR (DMSO- d_6 , δ) : 0.91 (3H, t, $J=7.1\text{Hz}$), 0.96 (3H, d, $J=6.7\text{Hz}$), 1.09 (3H, d, $J=5.7\text{Hz}$), 1.3-1.5 (4H, m),
25 1.7-2.6 (9H, m), 3.1-3.3 (1H, m), 3.7-4.6 (16H, m), 4.7-5.1 (7H, m), 5.11 (1H, d, $J=5.6\text{Hz}$), 5.17 (1H, d, $J=3.1\text{Hz}$), 5.26 (1H, d, $J=4.5\text{Hz}$), 5.55 (1H, d, $J=5.8\text{Hz}$), 6.7-6.9 (3H, m), 7.0-7.6 (6H, m), 7.97 (2H, d, $J=8.8\text{Hz}$), 8.0-8.4 (6H, m), 8.85 (1H, s),
30 8.92 (1H, d, $J=7.0\text{Hz}$)
FAB-MASS : $m/z = 1331$ (M+Na⁺)
Elemental Analysis Calcd. for C₅₅H₆₉N₁₀O₂₂NaS₂ :
C 45.45, H 5.89, N 9.64
Found : C 45.71, H 5.68, N 9.60

CLAIMS

1. A polypeptide compound of the following general formula :



[I]

wherein R^1 is lower alkanoyl substituted with

- unsaturated 6-membered heteromonocyclic group containing at least one nitrogen atom which may have one or more suitable substituent(s);
- lower alkanoyl substituted with 1,2,3,4-tetrahydroisoquinoline which may have one or more suitable substituent(s);
- lower alkanoyl substituted with unsaturated condensed heterocyclic group containing at least one oxygen atom which may have one or more suitable substituent(s);
- lower alkanoyl substituted with unsaturated condensed heterocyclic group containing 1 to 3 sulfur atom(s)

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aroyl substituted with aryl having heterocyclic(higher)alkoxy, in which heterocyclic group may have one or more suitable substituent(s);

aroyle substituted with aryl having
lower alkoxy(lower)alkoxy(higher)-
alkoxy;

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aroyle substituted with aryl having
lower alkoxy(lower)alkoxy;

aroyle substituted with aryl which has
aryl having lower alkoxy;

10

aroyle substituted with aryl which has
aryl having lower alkoxy(lower)alkoxy;

aroyle substituted with aryl having
heterocyclicoxy(higher)alkoxy;

aroyle substituted with aryl having
aryloxy(lower)alkoxy;

15

aroyle substituted with aryl having
heterocycliccarbonyl(higher)alkoxy;

lower alkanoyl substituted with
oxazolyl which has aryl having higher
alkoxy;

20

lower alkanoyl substituted with furyl
which has aryl substituted with aryl
having lower alkoxy;

lower alkanoyl substituted with
triazolyl which has oxo and aryl having
higher alkyl;

25

higher alkanoyl having hydroxy;

higher alkanoyl having ar(lower)alkyl
and hydroxy;

3-methyl-tridecenoyl; or

30

(C₂-C₆)alkanoyl substituted with aryl
having higher alkoxy, in which (C₂-C₆)-
alkanoyl may have amino or protected
amino, and

a pharmaceutically acceptable salt thereof.

35

2. A compound of claim 1, wherein

R¹ is lower alkanoyl substituted with unsaturated
6-membered heteromonocyclic group containing at
least one nitrogen atom which may have 1 to 3
substituent(s) selected from the group consisting
of lower alkoxy, higher alkoxy, lower alkyl,
higher alkyl, higher alkoxy(lower)alkyl, phenyl
having lower alkoxy, phenyl having higher alkoxy,
naphthyl having lower alkoxy, naphthyl having
higher alkoxy, phenyl having lower alkyl, phenyl
having higher alkyl, naphthoyl having higher
alkoxy, phenyl substituted with phenyl having
lower alkyl, 3 to 8-membered saturated
heteromonocyclic group containing at least one
nitrogen atom which may have phenyl having higher
alkoxy, phenyl substituted with phenyl having
lower alkoxy, 3 to 8-membered saturated
heteromonocyclic group containing at least one
nitrogen atom which may have phenyl having lower
alkoxy(higher)alkoxy, 3 to 8-membered saturated
heteromonocyclic group containing at least one
nitrogen atom which may have phenyl having lower
alkoxy, and oxo;

lower alkanoyl substituted with 1,2,3,4-
tetrahydroisoquinoline having higher alkoxy and
lower alkoxy carbonyl;

lower alkanoyl substituted with unsaturated
condensed heterocyclic group containing at least
one oxygen atom which may have 1 to 3
substituent(s) selected from the group consisting
of lower alkoxy, higher alkoxy, lower alkyl,
higher alkyl, higher alkoxy(lower)alkyl, phenyl
having lower alkoxy, phenyl having higher alkoxy,

naphthyl having lower alkoxy, naphthyl having higher alkoxy, phenyl having lower alkyl, phenyl having higher alkyl, naphthoyl having higher alkoxy, phenyl substituted with phenyl having lower alkyl, unsaturated 6-membered heteromonocyclic group containing at least one nitrogen atom which may have higher alkoxy, and oxo;

lower alkanoyl substituted with unsaturated condensed heterocyclic group containing 1 to 3 sulfur atom(s) which may have 1 to 3 substituent(s) selected from the group consisting of lower alkoxy, higher alkoxy, lower alkyl, higher alkyl, higher alkoxy(lower)alkyl, phenyl having lower alkoxy, phenyl having higher alkoxy, naphthyl having lower alkoxy, naphthyl having higher alkoxy, phenyl having lower alkyl, phenyl having higher alkyl, naphthoyl having higher alkoxy, phenyl substituted with phenyl having lower alkyl, and oxo;

lower alkanoyl substituted with unsaturated condensed heterocyclic group containing 2 or more nitrogen atoms which may have 1 to 3 substituent(s) selected from the group containing of lower alkoxy, higher alkoxy, lower alkyl, higher alkyl, higher alkoxy(lower)alkyl, phenyl having lower alkoxy, phenyl having higher alkoxy, naphthyl having lower alkoxy, naphthyl having higher alkoxy, phenyl having lower alkyl, phenyl having higher alkyl, naphthoyl having higher alkoxy, phenyl substituted with phenyl having lower alkyl, and oxo; or

lower alkanoyl substituted with saturated 3 to 8-membered heteromonocyclic group containing at least one nitrogen atom which may have 1 to 3

substituent(s) selected from the group consisting of lower alkoxy, higher alkoxy, lower alkyl, higher alkyl, higher alkoxy(lower)alkyl, phenyl having lower alkoxy, phenyl having higher alkoxy, naphthyl having lower alkoxy, naphthyl having higher alkoxy, phenyl having lower alkyl, phenyl having higher alkyl, naphthoyl having higher alkoxy, phenyl substituted with phenyl having lower alkyl, and oxo.

3. A compound of claim 1, wherein

R^1 is ar(lower)alkenoyl substituted with aryl which may have 1 to 3 substituent(s) selected from the group consisting of lower alkoxy, higher alkoxy, lower alkyl, higher alkyl, higher alkoxy(lower)alkyl, phenyl having lower alkoxy, phenyl having higher alkoxy, naphthyl having lower alkoxy, naphthyl having higher alkoxy, phenyl having lower alkyl, phenyl having higher alkyl, naphthoyl having higher alkoxy, phenyl substituted with phenyl having lower alkyl, lower alkoxy(lower)alkyl, halo(lower)alkoxy, lower alkenyloxy, halo(higher)alkoxy, lower alkoxy(higher)alkoxy, and oxo;

naphthyl(lower)alkenoyl which may have 1 to 3 higher alkoxy;

lower alkynoyl which may have 1 to 3 substituent(s) selected from the group consisting of lower alkoxy, higher alkoxy, lower alkyl, higher alkyl, higher alkoxy(lower)alkyl, phenyl having lower alkoxy, phenyl having higher alkoxy, naphthyl having lower alkoxy, naphthyl having higher alkoxy, phenyl having lower alkyl, phenyl having higher alkyl, naphthoyl having higher alkoxy, phenyl substituted with phenyl having

lower alkyl, and oxo;

ar(C₂-C₆)alkanoyl substituted with aryl having 1 to 3 substituent(s) selected from the group consisting of lower alkoxy, higher alkoxy, lower alkyl, higher alkyl, higher alkoxy(lower)alkyl, phenyl having lower alkoxy, phenyl having higher alkoxy, naphthyl having lower alkoxy, naphthyl having higher alkoxy, phenyl having lower alkyl, phenyl having higher alkyl, naphthoyl having higher alkoxy, phenyl substituted with phenyl having lower alkyl, phenyl having lower alkoxy(lower)alkoxy, and oxo, in which ar(C₂-C₆)-alkanoyl may have hydroxy, oxo, protected amino or amino; or

(C₂-C₆)alkanoyl substituted with naphthyl having higher alkoxy.

4. A compound of claim 1, wherein

R¹ is aroyl substituted with heterocyclic group which may have 1 to 3 substituent(s) selected from the group consisting of lower alkoxy, higher alkoxy, lower alkyl, higher alkyl, higher alkoxy(lower)alkyl, phenyl having lower alkoxy, phenyl having higher alkoxy, naphthyl having lower alkoxy, naphthyl having higher alkoxy, phenyl having lower alkyl, phenyl having higher alkyl, naphthoyl having higher alkoxy, phenyl substituted with phenyl having lower alkyl, phenyl having lower alkoxy(higher)alkoxy, phenyl having higher alkenyloxy, heterocyclic group substituted with phenyl having lower alkoxy, heterocyclic group, cyclo(lower)alkyl having phenyl, phenyl having cyclo(lower)alkyl, phenyl substituted with heterocyclic group having lower alkyl and oxo, cyclo(lower)alkyl having lower alkyl, phenyl

substituted with phenyl having lower alkoxy,
phenyl having heterocyclic group and oxo, in which
aroyl may have halogen;

5 aroyl substituted with aryl having
heterocyclic(higher)alkoxy, in which heterocyclic
group may have lower alkyl;

aroyl substituted with aryl having lower
alkoxy(higher)alkoxy;

10 aroyl substituted with aryl having lower
alkenyl(lower)alkoxy;

aroyl substituted with 2 lower alkoxy;

aroyl substituted with aryl having lower alkyl;
or

15 aroyl substituted with aryl having higher alkyl.

5. A compound of claim 1, wherein

20 R^1 is aryloxy(lower)alkanoyl which may have 1 to 3
substituent(s) selected from the group consisting
of lower alkoxy, higher alkoxy, lower alkyl,
higher alkyl, higher alkoxy(lower)alkyl, phenyl
having lower alkoxy, phenyl having higher alkoxy,
naphthyl having lower alkoxy, naphthyl having
higher alkoxy, phenyl having lower alkyl, phenyl
having higher alkyl, naphthoyl having higher
25 alkoxy, phenyl substituted with phenyl having
lower alkyl, and oxo;

30 ar(lower)alkoxy(lower)alkanoyl which may have 1
to 3 substituent(s) selected from the group
consisting of lower alkoxy, higher alkoxy, lower
alkyl, higher alkyl, higher alkoxy(lower)alkyl,
phenyl having lower alkoxy, phenyl having higher
alkoxy, naphthyl having lower alkoxy, naphthyl
having higher alkoxy, phenyl having lower alkyl,
phenyl having higher alkyl, naphthoyl having
35 higher alkoxy, phenyl substituted with phenyl

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having lower alkyl, and oxo; or

arylamino(lower)alkanoyl which may have 1 to 3
substituent(s) selected from the group consisting
of lower alkoxy, higher alkoxy, lower alkyl,
higher alkyl, higher alkoxy(lower)alkyl, phenyl
having lower alkoxy, phenyl having higher alkoxy,
naphthyl having lower alkoxy, naphthyl having
higher alkoxy, phenyl having lower alkyl, phenyl
having higher alkyl, naphthoyl having higher
alkoxy, phenyl substituted with phenyl having
lower alkyl, and oxo.

6. A compound of claim 1, wherein

R^1 is lower alkanoyl substituted with pyrazolyl which
has lower alkyl and aryl having higher alkoxy;

lower alkoxy(higher)alkanoyl, in which higher
alkanoyl may have amino or protected amino;

aroyl substituted with aryl having
heterocyclicoxy, in which heterocyclicoxy may have
phenyl;

aroyl substituted with cyclo(lower)alkyl having
lower alkyl;

indolylcarbonyl having higher alkyl;

naphthoyl having lower alkyl;

naphthoyl having higher alkyl;

naphthoyl having lower alkoxy(higher)alkoxy;

aroyl substituted with aryl having lower
alkoxy(lower)alkoxy(higher)alkoxy;

aroyl substituted with aryl having lower
alkoxy(lower)alkoxy;

aroyl substituted with aryl which has phenyl
having lower alkoxy;

aroyl substituted with aryl which has phenyl
having lower alkoxy(lower)alkoxy;

aroyl substituted with aryl having

heterocyclicoxy(higher)alkoxy;

aryyl substituted with aryl having
phenoxy(lower)alkoxy;

5 aryl substituted with aryl having
heterocycliccarbonyl(higher)alkoxy;

lower alkanoyl substituted with oxazolyl which
has aryl having higher alkoxy;

lower alkanoyl substituted with furyl which has
aryl substituted with phenyl having lower alkoxy;

10 lower alkanoyl substituted with triazolyl which
has oxo and phenyl having higher alkyl;

higher alkanoyl having hydroxy;

higher alkanoyl having benzyl and hydroxy;

3-methyl-tridecenoyl; or

15 (C₂-C₆)alkanoyl substituted with aryl having
higher alkoxy, in which (C₂-C₆)alkanoyl may have
amino or protected amino.

7. A compound of claim 2, wherein

20 R¹ is lower alkanoyl substituted with pyridyl or
pyridazinyl, each of which may have 1 to 3
substituent(s) selected from the group consisting
of higher alkoxy, higher alkoxy(lower)alkyl,
phenyl having higher alkoxy, phenyl substituted
25 with phenyl having lower alkoxy, piperazinyl
substituted with phenyl having higher alkoxy,
piperazinyl substituted with phenyl having lower
alkoxy(higher)alkoxy, and piperazinyl substituted
with phenyl having lower alkoxy;

30 lower alkanoyl substituted with 1,2,3,4-
tetrahydroisoquinoline having higher alkoxy and
lower alkoxy carbonyl;

lower alkanoyl substituted with coumarin which
may have 1 to 3 substituent(s) selected from the
35 group consisting of higher alkoxy, and oxo;

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lower alkanoyl substituted with benzothiophenyl
which may have 1 to 3 higher alkoxy;

lower alkanoyl substituted with benzo[b]furanyl
which may have 1 to 3 substituent(s) selected from
the group consisting of higher alkoxy and lower
alkyl;

lower alkanoyl substituted with benzoxazolyl
which may have 1 to 3 substituent(s) selected from
the group consisting of higher alkyl, phenyl
having lower alkoxy, phenyl substituted with
phenyl having lower alkyl, and pyridyl having
higher alkoxy;

lower alkanoyl substituted with benzimidazolyl
which may have 1 to 3 substituent(s) selected from
the group consisting of higher alkyl, and phenyl
having lower alkoxy; or

lower alkanoyl substituted with piperidyl or
piperazinyl, each of which may have 1 to 3
substituent(s) selected from the group consisting
of phenyl having higher alkoxy, and naphthoyl
having higher alkoxy.

8. A compound of claim 3, wherein

R^1 is phenyl(lower)alkenoyl substituted with phenyl
which may have 1 to 3 substituent(s) selected from
the group consisting of lower alkoxy, lower alkyl,
higher alkyl, lower alkoxy(lower)alkyl,
halo(lower)alkoxy, lower alkenyloxy,
halo(higher)alkoxy, and lower
alkoxy(higher)alkoxy;

naphthyl(lower)alkenoyl which may have 1 to 3
higher alkoxy;

lower alkynoyl which may have 1 to 3
substituent(s) selected from the group consisting
of naphthyl having higher alkoxy, and phenyl

substituted with phenyl having lower alkyl;

phenyl(C₂-C₆)alkanoyl substituted with phenyl
which has 1 to 3 substituent(s) selected from the
group consisting of lower alkoxy, higher alkoxy,
lower alkyl, higher alkyl, and phenyl having lower
alkoxy(lower)alkyl,

in which phenyl(C₂-C₆)alkanoyl may have hydroxy,
oxo, protected amino or amino; or

(C₂-C₆)alkanoyl substituted with naphthyl having
higher alkoxy.

9. A compound of claim 4, wherein

R¹ is benzoyl substituted with saturated 6-membered
heteromonocyclic group containing at least one
nitrogen atom which may have 1 to 3 substituent(s)
selected from the group consisting of phenyl
having lower alkoxy, phenyl having higher alkoxy,
phenyl having lower alkyl, phenyl having lower
alkoxy(higher)alkoxy, phenyl having higher
alkenyloxy, piperidyl substituted with phenyl
having lower alkoxy, piperidyl, cyclo(lower)alkyl
having phenyl, phenyl having cyclo(lower)alkyl,
and phenyl substituted with triazolyl having oxo
and lower alkyl,

in which benzoyl may have halogen;

benzoyl substituted with unsaturated 5-membered
heteromonocyclic group containing 1 to 2 oxygen
atom(s) and 1 to 3 nitrogen atom(s) which may have
1 to 3 substituent(s) selected from the group
consisting of higher alkyl, phenyl having lower
alkoxy, phenyl having higher alkoxy, phenyl having
lower alkoxy(higher)alkoxy, and phenyl substituted
with phenyl having lower alkoxy;

benzoyl substituted with 5 or 6-membered
heteromonocyclic group containing 1 or 2 nitrogen

atom(s) which may have 1 to 3 substituent(s) selected from the group consisting of higher alkyl and phenyl having lower alkoxy;

5 benzoyl substituted with 5-membered heteromonocyclic group containing 1 to 2 nitrogen atom(s) and 1 to 2 sulfur atom(s) which may have 1 to 3 substituent(s) selected from the group consisting of phenyl having lower alkoxy, phenyl having higher alkoxy, cyclo(lower)alkyl having
10 lower alkyl, phenyl substituted with phenyl having lower alkoxy, phenyl having cyclo(lower)alkyl, phenyl having piperidine, and phenyl having lower alkoxy(higher)alkoxy;

15 benzoyl substituted with phenyl having higher alkoxy substituted with unsaturated 3 to 8-membered heteromonocyclic group containing at least one nitrogen atom;

20 benzoyl substituted with phenyl having higher alkoxy substituted with saturated 6-membered heteromonocyclic group containing 1 to 2 oxygen atom(s) and 1 to 3 nitrogen atom(s) which may have lower alkyl;

25 benzoyl substituted with phenyl having lower alkoxy(higher)alkoxy;

benzoyl substituted with phenyl having lower alkenyl(lower)alkoxy;

benzoyl substituted with 2 lower alkoxy;

benzoyl substituted with phenyl having lower alkyl; or

30 benzoyl substituted with phenyl having higher alkyl.

10. A compound of claim 5, wherein

35 R^1 is phenyloxy(lower)alkanoyl which may have 1 to 3 higher alkoxy;

phenyl(lower)alkoxy(lower)alkanoyl which may have 1 to 3 higher alkoxy; or

phenylamino(lower)alkanoyl which may have 1 to 3 higher alkoxy.

5

11. A compound of claim 1, wherein

R¹ is benzoyl substituted with piperazinyl which may have 1 to 3 substituent(s) selected from the group consisting of phenyl having lower alkoxy, phenyl having higher alkoxy, phenyl having lower alkyl, phenyl having lower alkoxy(higher)alkoxy, phenyl having higher alkenyloxy, piperidyl substituted with phenyl having lower alkoxy, cyclo(lower)alkyl having phenyl, phenyl having cyclo(lower)alkyl, and phenyl substituted with triazolyl having oxo and lower alkyl, in which benzoyl may have halogen;

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benzoyl substituted with isoxazolyl which may have 1 to 3 substituent(s) selected from the group consisting of higher alkyl, phenyl having lower alkoxy, phenyl having higher alkoxy, phenyl having lower alkoxy(higher)alkoxy, and phenyl substituted with phenyl having lower alkoxy;

20

benzoyl substituted with phenyl having lower alkoxy(higher)alkoxy;

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benzoyl substituted with phenyl having lower alkyl;

benzoyl substituted with phenyl having higher alkyl;

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phenyl(lower)alkenoyl substituted with phenyl which may have 1 to 3 substituent(s) selected from the group consisting of lower alkoxy, lower alkyl, higher alkyl, lower alkoxy(lower)alkyl, halo(lower)alkoxy, lower alkenyloxy, halo(higher)alkoxy and lower alkoxy(higher)alkoxy;

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benzoyl substituted with thiadiazolyl which may have 1 to 3 substituent(s) selected from the group consisting of phenyl having lower alkoxy, phenyl having higher alkoxy, cyclo(lower)alkyl having lower alkyl, phenyl substituted with phenyl having lower alkoxy, phenyl having cyclo(lower)alkyl, phenyl having piperidyl, and phenyl having lower alkoxy(higher)alkoxy; or

benzoyl substituted with oxadiazolyl which may have 1 to 3 substituent(s) selected from the group consisting of phenyl having lower alkoxy, phenyl having higher alkoxy, phenyl having lower alkoxy(higher)alkoxy, higher alkyl and phenyl substituted with phenyl having lower alkoxy.

12. A compound of claim 11, wherein

R^1 is benzoyl substituted with phenyl having lower alkoxy(higher)alkoxy; or

benzoyl substituted with phenyl having lower alkyl.

13. A compound of claim 11, wherein

R^1 is benzoyl substituted with piperazinyl which may have phenyl having lower alkoxy;

benzoyl substituted with isoxazolyl which may have phenyl having lower alkoxy;

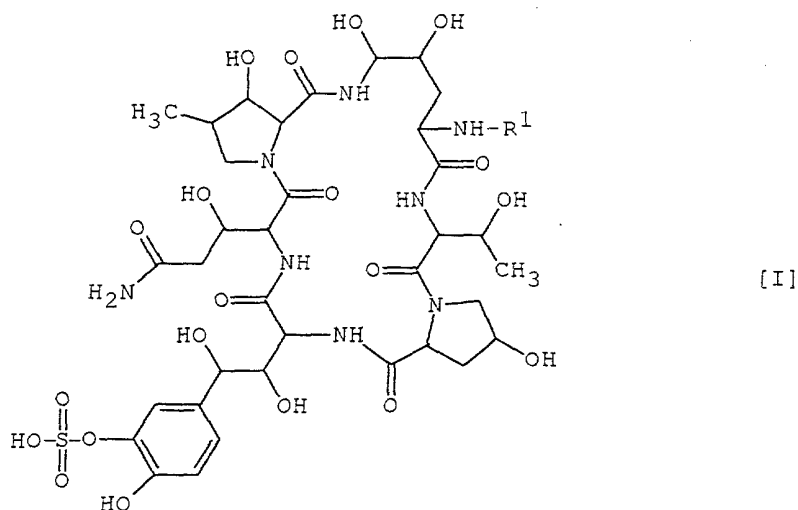
benzoyl substituted with thiadiazolyl which may have phenyl having lower alkoxy(higher)alkoxy; or

benzoyl substituted with oxadiazolyl which may have phenyl having lower alkoxy.

14. A compound of claim 11, wherein

R^1 is phenyl(lower)alkenoyl substituted with phenyl which may have lower alkoxy.

15. A process for the preparation of a polypeptide compound of the formula [I] :



wherein

R^1 is lower alkanoyl substituted with unsaturated 6-membered heteromonocyclic group containing at least one nitrogen atom which may have one or more suitable substituent(s);

lower alkanoyl substituted with 1,2,3,4-tetrahydro-isoquinoline having higher alkoxy;

lower alkanoyl substituted with unsaturated condensed heterocyclic group containing at least one oxygen atom which may have one or more suitable substituent(s);

lower alkanoyl substituted with unsaturated condensed heterocyclic group containing 1 to 3 sulfur atom(s) which may have one or more suitable substituent(s);

lower alkanoyl substituted with unsaturated condensed heterocyclic group containing 2 or more nitrogen atom(s) which may have one or more suitable substituent(s);

lower alkanoyl substituted with saturated 3 to 8-membered heteromonocyclic group containing at least one nitrogen atom which may have one or more suitable substituent(s);

5 ar(lower)alkenoyl substituted with aryl which may have one or more suitable substituent(s);

 naphthyl(lower)alkenoyl which may have one or more higher alkoxy;

10 lower alkynoyl which may have one or more suitable substituent(s);

 (C₂-C₆)alkanoyl substituted with naphthyl having higher alkoxy;

15 ar(C₂-C₆)alkanoyl substituted with aryl having one or more suitable substituent(s), in which ar(C₂-C₆)alkanoyl may have one or more suitable substituent(s);

20 aroyl substituted with heterocyclic group which may have one or more suitable substituent(s), in which aroyl may have one or more suitable substituent(s);

 aroyl substituted with aryl having heterocyclic(higher)alkoxy, in which heterocyclic group may have one or more suitable substituent(s);

25 aroyl substituted with aryl having lower alkoxy(higher)alkoxy;

 aroyl substituted with aryl having lower alkenyl(lower)alkoxy;

 aroyl substituted with 2 lower alkoxy;

30 aroyl substituted with aryl having lower alkyl;

 aroyl substituted with aryl having higher alkyl;

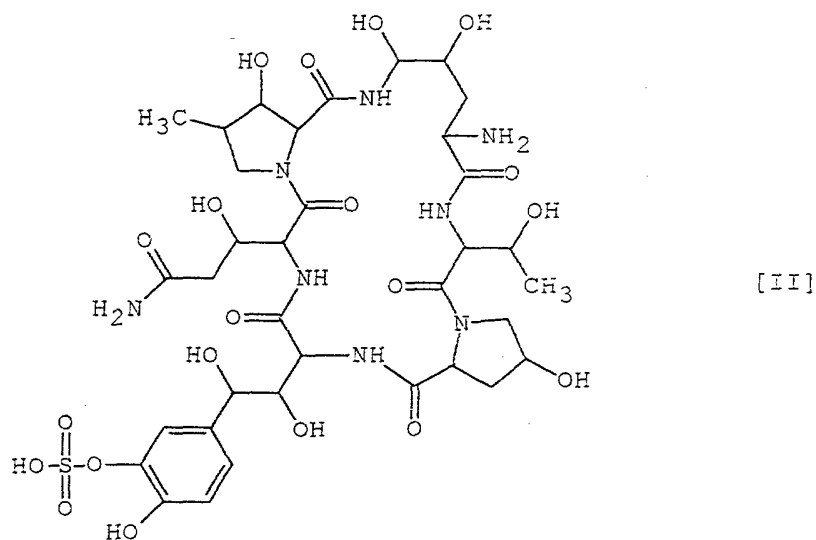
 aryloxy(lower)alkanoyl which may have one or more suitable substituent(s);

35 ar(lower)alkoxy(lower)alkanoyl which may have one or more suitable substituent(s);

has oxo and aryl having higher alkyl;
higher alkanoyl having hydroxy;
higher alkanoyl having ar(lower)alkyl and
hydroxy;

3-methyl-tridecenoyl; or
(C₂-C₆)alkanoyl substituted with aryl having
higher alkoxy, in which (C₂-C₆)alkanoyl may have
amino or protected amino, and
a pharmaceutically acceptable salt thereof,
which comprises

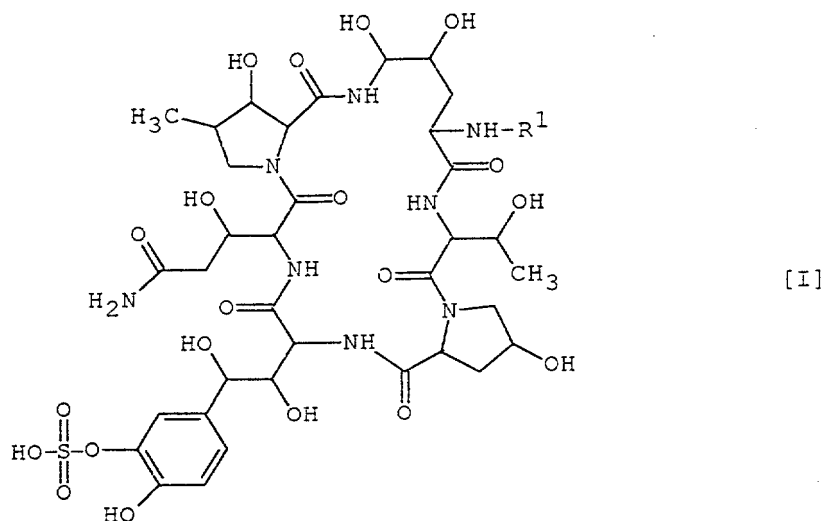
1) reacting a compound of the formula :



or its reactive derivative at the amino group or a salt
thereof, with a compound of the formula :



wherein R¹ is defined above,
or its reactive derivative at the carboxy group or a
salt thereof, to give a compound [I] of the formula :



wherein R¹ is defined above,
or a salt thereof.

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16. A pharmaceutical composition which comprises, as an active ingredient, a compound of claim 1 or a pharmaceutically acceptable salt thereof in admixture with pharmaceutically acceptable carriers or excipients.

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17. Use of a compound of claim 1 or a pharmaceutically acceptable salt thereof as a medicament.

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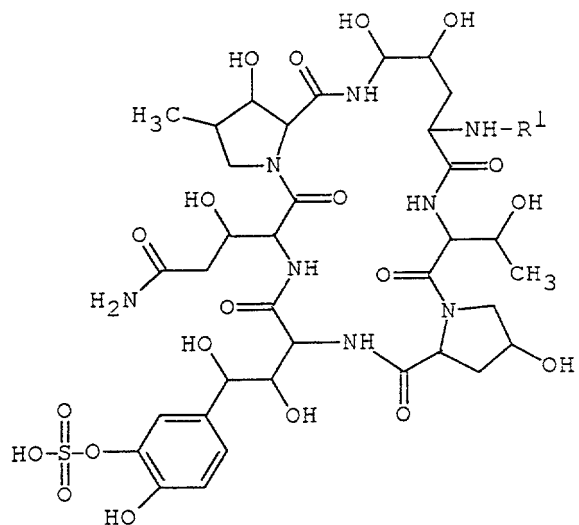
18. A compound of claim 1 or a pharmaceutically acceptable salt thereof for use as a medicament.

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19. A method for the prophylactic and/or the therapeutic treatment of infectious diseases caused by pathogenic microorganisms which comprises administering a compound of claim 1 or a pharmaceutically acceptable salt thereof to a human being or an animal.

ABSTRACT

This invention relates to new polypeptide compounds represented by the following formula (I) :



wherein

R^1 is as defined in the description and pharmaceutically acceptable salt thereof which have antimicrobial activities (especially, antifungal activities), inhibitory activity on β -1,3-glucan synthase, to process for preparation thereof, to a pharmaceutical composition comprising the same, and to a method for the prophylactic and/or therapeutic treatment of infectious diseases including Pneumocystis carinii infection (e.g. Pneumocystis carinii pneumonia) in a human being or an animal.

Declaration, Power Of Attorney and Petition

Page 1 of 3

WE (I) the undersigned inventor(s), hereby declare(s) that:

My residence, post office address and citizenship are as stated below next to my name,

We (I) believe that we are (I am) the original, first, and joint (sole) inventor(s) of the subject matter which is claimed and for which a patent is sought on the invention entitled

CYCLIC HEXAPEPTIDES HAVING ANTIBIOTIC ACTIVITY

the specification of which

- ☐ is attached hereto.
- ☐ was filed on _____ as
Application Serial No. _____
and amended on _____.
- ☒ was filed as PCT international application
Number _____ PCT/JP95/01983
on _____ September 29, 1995,
and was amended under PCT Article 19
on _____ (if applicable).

We (I) hereby state that we (I) have reviewed and understand the contents of the above-identified specification, including the claims, as amended by any amendment referred to above.

We (I) acknowledge the duty to disclose information known to be material to the patentability of this application as defined in Section 1.56 of Title 37 Code of Federal Regulations.

We (I) hereby claim foreign priority benefits under 35 U.S.C. § 119(a)-(d) or § 365(b) of any foreign application(s) for patent or inventor's certificate, or § 365(a) of any PCT International application which designated at least one country other than the United States, listed below and have also identified below, by checking the box, any foreign application for patent or inventor's certificate, or PCT International application having a filing date before that of the application on which priority is claimed. Prior Foreign Application(s)

Application No.	Country	Day/Month/Year	Priority Claimed
9420425.2	G. Britain	07/10/94	<input checked="" type="checkbox"/> Yes <input type="checkbox"/> No
9508745.8	G. Britain	28/04/95	<input checked="" type="checkbox"/> Yes <input type="checkbox"/> No
_____	_____	_____	<input type="checkbox"/> Yes <input type="checkbox"/> No
_____	_____	_____	<input type="checkbox"/> Yes <input type="checkbox"/> No

We (I) hereby claim the benefit under Title 35, United States Code, § 119(e) of any United States provisional application(s) listed below.

_____	_____
(Application Number)	(Filing Date)
_____	_____
(Application Number)	(Filing Date)

We (I) hereby claim the benefit under 35 U.S.C. § 120 of any United States application(s), or § 365(c) of any PCT International application designating the United States, listed below and, insofar as the subject matter of each of the claims of this application is not disclosed in the prior United States or PCT International application in the manner provided by the first paragraph of 35 U.S.C. § 112, I acknowledge the duty to disclose information which is material to patentability as defined in 37 CFR § 1.56 which became available between the filing date of the prior application and the national or PCT International filing date of this application.

Application Serial No.	Filing Date	Status (pending, patented, abandoned)
PCT/JP95/01983	September 29, 1995	_____
_____	_____	_____
_____	_____	_____

And we (I) hereby appoint: Norman F. Oblon, Registration Number 24,618; Marvin J. Spivak, Registration Number 24,913; C. Irvin McClelland, Registration Number 21,124; Gregory J. Maier, Registration Number 25,599; Arthur I. Neustadt, Registration Number 24,854; Richard D. Kelly, Registration Number 27,757; James D. Hamilton, Registration Number 28,421; Eckhard H. Kuesters, Registration Number 28,870; Robert T. Pous, Registration Number 29,099; Charles L. Gholz, Registration Number 26,395; Vincent J. Sunderdick, Registration Number 29,004; William E. Beaumont, Registration Number 30,996; Steven B. Kelber, Registration Number 30,073; Robert F. Gnuse, Registration Number 27,295; Jean-Paul Lavalleye, Registration Number 31,451; Timothy R. Schwartz, Registration Number 32,171; Stephen G. Baxter, Registration Number 32,884; Martin M. Zoltick, Registration Number 35,745; Robert W. Hahl, Registration Number 33,893; Richard L. Treanor, Registration Number 36,379; Steven P. Weihrouch, Registration Number 32,829; John T. Goolkasian, Registration Number 26,142; Marc R. Labgold, Registration Number 34,651; William J. Healey, Registration Number 36,160; Richard L. Chinn, Registration Number 34,305; Steven E. Lipman, Registration Number 30,011; and Jacques M. Dulin, Registration Number 24,067; our (my) attorneys, with full powers of substitution and revocation, to prosecute this application and to transact all business in the Patent Office connected therewith; and we (I) hereby request that all correspondence regarding this application be sent to the firm of OBLON, SPIVAK, McCLELLAND, MAIER & NEUSTADT, P.C., whose Post Office Address is: Fourth Floor, 1755 Jefferson Davis Highway, Arlington, Virginia 22202.

We (I) declare that all statements made herein of our (my) own knowledge are true and that all statements made on information and belief are believed to be true; and further that these statements were made with the knowledge that willful false statements and the like so made are punishable by fine or imprisonment, or both, under Section 1001 of Title 18 of the United States Code and that such willful false statements may jeopardize the validity of the application or any patent issuing thereon.

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